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(54) Title: PULMONARY SURFACTANT PROTEIN AND RELATED POLYPEPTIDE

(57) Abstract

The present invention relates to a human SP18 monomer protein-related polypeptide useful in forming a synthetic pulmonary surfactant. The present invention also relates to a method of treating neonatal respiratory distress syndrome comprising administering a therapeutically effective amount of synthetic pulmonary of the present invention. Further contemplated by the present invention is a composition containing human SP18 monomer and human SP18 dimer but no other pulmonary surfactant proteins. A recombinant DNA molecule capable of expressing, without post-translational proteolytic processing, mature human SP18 monomer, and methods of using the recombinant DNA molecule are also contemplated.

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PULMONARY SURFACTANT PROTEIN
AND RELATED POLYPEPTIDE

Description

5 Cross Reference to Related Application

This is a Continuation-In-Part application of copending application Serial Number 141,200, filed January 6, 1988, entitled Pulmonary Surfactant Protein and Related Polypeptide (Cochrane, et al), which 10 application is hereby incorporated by reference.

Technical Field

The present invention relates to SP18 monomer-related polypeptides useful in forming synthetic 15 pulmonary surfactants. The present invention also relates to a recombinant nucleic acid molecule carrying a structural gene that encodes human SP18 monomer protein and the use of such a recombinant molecule to produce human SP18 monomer.

20

Background

Pulmonary surfactant (PS) lines the alveolar epithelium of mature mammalian lungs. Natural PS has been described as a "lipoprotein complex" because it 25 contains both phospholipids and apoproteins that interact to reduce surface tension at the lung air-liquid interface.

Since the discovery of pulmonary surfactant, and the subsequent finding that its deficiency was the 30 primary cause of neonatal respiratory distress syndrome (RDS), a number of studies have been directed towards developing effective surfactant replacement therapy for affected individuals, particularly infants, using exogenous PS. For instance, 35 improvements in lung function as measured by a

decrease in mean airway pressure and oxygen requirements have been demonstrated using exogenous surfactants in human pre-term infants. See Hallman, et al, Pediatrics, 71:473-482 (1983); Merritt, et al,

5 5 J. Pediatr., 108:741-745 (1986); Hallman, et al, J. Pediatr., 106:963-969 (1985); Morley, et al, Lancet, i:64-68 (1981); Merritt, et al, New England J. Med., 315:785-790 (1986), Smyth, et al, Pediatrics, 71:913-917 (1983); Enhornig, et al, Pediatrics, 76:145-153

10 10 (1985); Fujiwara, et al, The Lancet, 1:55-59 (1980); Kwong, et al, Pediatrics, 76:585-592 (1985); Shapiro, et al, Pediatrics, 76:593-599 (1985); Fujiwara, in "Pulmonary Surfactant", Robertson, B., Van Golde, L.M.G., Batenburg J. (eds), Elsevier Science

15 15 Publishers, Amsterdam, pp. 479-503, (1984).

From a pharmacologic point of view, the optimal exogenous PS to use in the treatment of RDS would be one completely synthesized in the laboratory, under controlled and sterile conditions, with negligible 20 batch-to-batch variability in properties. To minimize the possibility of immunologic complications, the apoprotein component of an exogenous PS should be identical to that found in humans. Unfortunately, the composition of naturally occurring PS is complex, and 25 the art has not yet identified all of the biochemical components that generate the biophysical properties needed for high physiologic activity in lungs. In particular, the art has failed to characterize all of the apoproteins present in natural PS or identify the 30 function of the PS apoproteins presently known.

It should be noted that the literature on PS apoproteins and their roles in surfactant function is complex, inconsistent and sometimes contradictory because heterogenous apoprotein preparations were used 35 in many studies. To date, the art has not

definitively established the number of different apoproteins present in natural PS.

Of particular interest to the present invention is the use of a low molecular weight (LMW) human PS-

5 associated apoprotein as a component in an exogenous surfactant. Several studies have attempted to isolated or define human PS LMW apoproteins using biochemical techniques. See, for example,
Phizackerley, et al, Biochem. J., 183:731-736 (1979),
10 Revak, et al, Am. Rev. Resp. Dis., 134:1258-1265 (1986), Suzuki, et al, Eur. J. Respir. Dis., 69:335-345 (1986), Taeusch, et al, Pediatrics, 77:572-581 (1986), Yu, et al, Biochem. J., 236:85-89 (1986), Whitsett, et al, Pediatric Res., 20:460-467 (1986),
15 Whitsett, et al, Pediatric Res., 20:744-749 (1986), Takahashi, et al, Biochem. Biophys. Res. Comm., 135:527-532 (1986), Suzuki, et al, Exp. Lung. Res., 11:61-73 (1986), Curstedt, et al, Eur. J. Biochem., 168:255-262 (1987), Notter, et al, Chem. Phys. Lipids,
20 44:1-17 (1987), and Phelps, et al, Am. Rev. Respir. Dis., 135:1112-1117 (1987).

Recently, the art has begun to apply the methods of recombinant DNA technology to overcome the problems associated with not being able to isolate to

25 homogeneity the individual LMW PS apoproteins. For instance, Glasser, et al, Proc. Natl. Acad. Sci., U.S.A., 84:4007-4011 (1987) reported a cDNA derived sequence of amino acid residues that forms at least a portion of a human precursor protein from which at
30 least one mature LMW apoprotein, which they designated SPL (Phe), is formed. While Glasser, et al were not able to determine the carboxy-terminal residue of SPL(Phe), and therefore were not able to identify its complete sequence, they did predict that mature
35 SPL(Phe) was about 60 amino acids in length.

Jacobs, et al, J. Biol. Chem., 262:9808-9811 (1987) have described a cDNA and derived amino acid residue sequence for a human precursor protein similar to that described by Glasser, et al, supra. However, 5 according to Jacobs et al. the mature LMW apoprotein, which they designated PSP-B, formed from the precursor would be 76 amino acid residues in length. In addition, Jacobs, et al, noted that it was not clear that any PS apoprotein derived from the reported 10 precursor protein was present in the surfactant preparations that had been studied clinically by others.

From the foregoing it can be seen that the literature contains multiple nomenclature for what is 15 apparently the same PS apoprotein. Therefore, for ease of discussion, the mature apoprotein derived from the precursor protein described by Glasser, et al, supra, and Jacobs, et al, supra, will be referred to herein generically as "SP18", with the monomeric and 20 dimeric forms being referred to as "SP18 monomer" and "SP18 dimer", respectively, when appropriate.

The canine SP18 precursor has been described by Hawgood, et al, Proc. Natl. Acad. Sci. U.S.A., 84:66-70 (1987) and Schilling, et al, International Patent 25 Application WO 86/03408. However, it should be noted that both those studies suffered the same inability to define the mature, biologically active form of SP18 as the Glasser, et al, supra, and Jacobs, et al, supra, studies.

30 Warr, et al, Proc. Natl. Acad. Sci., U.S.A., 84:7915-7919 (1987) describe a cDNA derived sequence of 197 amino acid residues that forms a precursor protein from which a mature LMW apoprotein, they designate as SP5, is formed. Like the studies 35 attempting to describe SP18, Warr, et al, were unable

to determine the carboxy terminal residue of the mature protein formed from the precursor protein sequence, and thus were not able to definitively characterize SP5.

5 Because the amino acid residue sequence of the precursor protein reported by Warr, et al, is different from that reported by Glasser, et al, and Jacobs, et al, it therefore appears that the art has determined that natural PS contains at least two LMW 10 apoproteins. However, the biologically active forms of those proteins has remained undetermined.

Summary

It has now been found that human SP18 is a 15 homodimeric protein (SP18 dimer) whose mature subunit protein (SP18 monomer) displays an apparent molecular weight of about 9,000 daltons when determined by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE).

20 It has also been discovered that human SP18 can function as an active ingredient in a synthetic pulmonary surfactant in the absence of other previously identified pulmonary surfactant proteins.

25 In addition, the carboxy-terminal amino acid residue sequence of the mature human SP18 monomer protein has been determined, and thus its naturally occurring form has now been discovered.

30 Thus, the present invention contemplates a composition comprising substantially isolated or substantially pure human SP18 monomer.

Also contemplated by the present invention is a DNA segment consisting essentially of a DNA sequence encoding human SP18 monomer protein.

35 Another embodiment of the present invention is a recombinant nucleic acid molecule comprising a vector

operatively linked to a structural gene capable of expressing, without post-translational proteolytic modification, human SP18 monomer protein.

Also contemplated is a method of preparing human 5 SP18 monomer protein comprising:

(a) initiating a culture, in a nutrient medium, of mammalian cells transformed with a recombinant DNA molecule comprising a vector operatively linked to a structural gene capable of expressing, without post-10 translational proteolytic modification, human SP18 monomer;

(b) maintaining said culture for a time period sufficient for said cells to express human SP18 monomer protein from said recombinant DNA molecule; 15 and

(c) recovering said protein from said culture.

Further contemplated is a polypeptide consisting essentially of at least 10 amino acid residues and no more than about 60 amino acid residues corresponding 20 in sequence to the amino acid residue sequence of human SP18 monomer, said polypeptide, when admixed with a pharmaceutically acceptable phospholipid form a synthetic surfactant having a surfactant activity greater than the surfactant activity of the 25 phospholipid alone.

In another embodiment, the subject invention contemplates a synthetic pulmonary surfactant comprising a pharmaceutically acceptable phospholipid admixed with a polypeptide consisting essentially of 30 at least 10 amino acid residues and no more than about 60 amino acid residues corresponding in sequence to the amino acid residue sequence of human SP18 monomer.

A further embodiment of the subject invention is a method of treating respiratory distress syndrome 35 comprising administering a therapeutically effective

amount of a synthetic pulmonary surfactant, said surfactant comprising a pharmaceutically acceptable phospholipid admixed with an effective amount of a polypeptide consisting essentially of at least 10 5 amino acid residues and no more than about 60 amino acid residues corresponding in sequence to the amino acid residue sequence of human SP18 monomer, said polypeptide, when admixed with a pharmaceutically acceptable phospholipid forms a synthetic surfactant 10 having a surfactant activity greater than the surfactant activity of the phospholipid alone.

Another embodiment of the subject invention is a method of treating respiratory distress syndrome comprising administering a therapeutically effective 15 amount of a synthetic pulmonary surfactant, said surfactant comprising an effective amount of either substantially isolated human SP18 monomer or substantially pure human SP18 monomer admixed with a pharmaceutically acceptable phospholipid.

20

Brief Description of Drawings

Figure 1 illustrates a 750 nucleotide cDNA sequence (top lines) and deduced amino acid residue sequence (bottom lines). The number to the right of 25 each line of nucleotides represents the numerical position in the sequence of the nucleotide at the end of each line. The nucleotides are grouped into codons, 15 codons per line, with the amino acid residue coded for by each codon shown in triple letter 30 code directly below the codon. The numerical position of some residues in the amino acid residue sequence encoded by the cDNA is shown below the residues. The amino-terminal amino acid residue of mature human SP18 monomer is Phe (encoded by nucleotides 187-189) and is 35 designated residues number 1. The carboxy-terminal

amino acid residue is Asp at residue position 81 (encoded by nucleotides 427-429). A structural gene encoding mature SP18 monomer therefore contains 81 codons and has a nucleotide sequence that corresponds 5 to nucleotides 187-429.

Figure 2 illustrates the protein elution profile of PS apoproteins from a Bio-Sil HA (silicic acid) column. Results of Pierce BCA protein assay (solid line) and phospholipid analyses (broken line) are 10 shown for selected fractions. Two milliliters (ml) were collected per fraction. Positive protein assay in fractions 28 to 33 is due to the presence of phospholipids.

Figure 3 illustrates a Silver-stained SDS-PAGE of 15 low molecular weight (LMW) PS apoproteins. Lanes A and D show a sample after silicic acid or Sephadex LH-20 chromatography; both LMW proteins are present. Lanes B, C, E and F show the resolution of SP18 (lanes 20 B and E) and SP9 (lanes C and F) following chromatography on Sephadex LH-60. Molecular weight standards are shown in lane G. Lanes A-C are unreduced samples, lanes D-F contain identical samples reduced with β -mercaptoethanol prior to electrophoresis.

Figure 4 illustrates inflation and deflation 25 pressure/volume curves of fetal rabbit lungs 30 min after intratracheal instillation of 100 μ l of saline (open circles), 2 mg phospholipids (PL) DPPC:PG, 3:1 (closed circles), PL + 10 μ g SP9 (open squares) PL + 10 μ g SP18 (closed squares), or 2 mg natural human 30 surfactant (closed triangles). Data are expressed as the mean of 4 animals \pm one standard deviation.

Figure 5 illustrates fetal rabbit lung tissue samples (x125 magnification, hematoxylin-eosin stain) following treatment with saline (A), natural human

surfactant (B), phospholipids DPPC:PG (C) or phospholipids plus LMW apoproteins (SP9 + SP18) (D).

Figure 6 illustrates the surfactant activity of exemplary polypeptide containing synthetic surfactants 5 of the present invention. Surfactant activity was determined by measuring the pressure gradient across an air/liquid interface using the pulsating bubble technique. The pressure gradient (ΔP) across the surface of the bubble is the absolute value of the 10 pressure recorded in centimeters of H_2O . The results obtained for each synthetic pulmonary surfactant are identified by the polypeptide in the surfactant. Results obtain for surfactants consisting of phospholipid alone (i.e., with no peptide or protein 15 admixed therewith) are identified as PL. The results obtained using a control peptide having only 8 amino acid residues and having a sequence corresponding to human SP18 monomer residues 74-81 (p74-81) are also shown. The data time points shown were obtained at 15 20 seconds, 1 minute and 5 minutes.

Figure 7 is a series of two graphs that illustrate the results of a static compliance study of exemplary synthetic surfactants of this invention using the fetal rabbit model previously described in Revak, et 25 al, Am. Rev. Respir. Dis., 134:1258-1265 (1986).

Following instillation of a synthetic surfactant or control into the trachea, the rabbit was ventilated for 30 minutes prior to making static compliance measurements. The "x" axis represents the pressure in 30 cm of water, while the "y" axis represents the volume in ml/kg of body weight. The graph on the left represents values at inflation and the graph on the right represents deflation values. The results for the following tested surfactants are illustrated: 35 natural surfactant (open square with a dot in the

center), phospholipid (PL) with 7% p52-81 (a polypeptide corresponding to residues 52 to 81 of SP18) (closed diamonds); PL with 3% P52-81 (closed squares with white dot in center); PL with 7% p36-81 5 (open diamonds); PL with 3% p66-81 (closed squares); PL with 3% p1-15 (open squares) and PL control (closed triangles).

Figure 8 is a series of two graphs that illustrate the results of a static compliance study of exemplary 10 synthetic surfactants of the invention. The procedure was performed as described in Figure 7 except that a different instillation procedure was used. The "x" and "y" axis and right and left graphs are as described in Figure 7. The results for the following 15 tested surfactants are illustrated: natural surfactant (open squares with a dot in the center); phospholipid (PL) with 10% p51-81 (closed diamonds); PL with 10% p51-76 (closed squares); and PL (closed triangles).

20

Detailed Description of the Invention

A. Definitions

Amino Acid: All amino acid residues identified herein are in the natural L-configuration. 25 In keeping with standard polypeptide nomenclature, J. Biol. Chem., 243:3557-59, (1969), abbreviations for amino acid residues are as shown in the following Table of Correspondence:

TABLE OF CORRESPONDENCE

SYMBOL			
	<u>1-Letter</u>	<u>3-Letter</u>	<u>AMINO ACID</u>
5	Y	Tyr	L-tyrosine
	G	Gly	glycine
	F	Phe	L-phenylalanine
	M	Met	L-methionine
	A	Ala	L-alanine
	S	Ser	L-serine
10	I	Ile	L-isoleucine
	L	Leu	L-leucine
	T	Thr	L-threonine
	V	Val	L-valine
	P	Pro	L-proline
	K	Lys	L-lysine
15	H	His	L-histidine
	Q	Gln	L-glutamine
	E	Glu	L-glutamic acid
	W	Trp	L-tryptophan
	R	Arg	L-arginine
	D	Asp	L-aspartic acid
20	N	Asn	L-asparagine
	C	Cys	L-cysteine

25 It should be noted that all amino acid residue sequences are represented herein by formulae whose left to right orientation is in the conventional direction of amino-terminus to carboxy-terminus. Furthermore, it should be noted that a dash at the

30 beginning or end of an amino acid residue sequence indicates a bond to a radical such as H and OH (hydrogen and hydroxyl) at the amino- and carboxy-termini, respectively, or a further sequence of one or more amino acid residues. In addition, it should be

35 noted that a virgule (/) at the right-hand end of a

other alveolar surfactant proteins and wherein less than 20 percent, preferably less than 10 percent and more preferably less than 5 percent, of the SP18 monomer present is in homodimeric form, i.e., present 5 as part of SP18 dimer.

Preferably, a SP18 monomer-containing composition of the present invention contains human SP18 monomer. More preferably, a SP18 monomer-containing composition contains SP18 monomer having an 10 amino acid residue sequence corresponding to the amino acid residue sequence shown in Figure 1 from about residue position 1 to at least about residue position 75, preferably to at least about position 81. More preferably, a SP18 monomer used to form a subject 15 protein composition corresponds in sequence to the sequence shown in Figure 1 from residue position 1 to residue position 81.

Preferably, the amino acid residue sequence of a SP18 monomer in a subject SP18 monomer-containing 20 composition corresponds to the sequence of a native SP18 monomer. However, it should be understood that a SP18 monomer used to form a protein composition of the present invention need not be identical to the amino acid residue sequence of a native SP18 monomer, but 25 may be subject to various changes, such as those described hereinbelow for a polypeptide of this invention, so long as such modifications do not destroy surfactant activity. Such modified protein can be produced, as is well known in the art, through, 30 for example, genomic site-directed mutagenesis.

"Surfactant activity" for a protein or polypeptide is defined as the ability, when combined with lipids, either alone or in combination with other proteins, to exhibit activity in the in vivo assay of 35 Robertson, Lung, 158:57-68 (1980). In this assay, the

residue sequence indicates that the sequence is continued on the next line.

Polypeptide and Peptide: Polypeptide and peptide are terms used interchangeably herein to designate a 5 linear series of no more than about 60 amino acid residues connected one to the other by peptide bonds between the alpha-amino and carboxy groups of adjacent residues.

Protein: Protein is a term used herein to 10 designate a linear series of greater than about 60 amino acid residues connected one to the other as in a polypeptide.

Nucleotide: a monomeric unit of DNA or RNA consisting of a sugar moiety (pentose), a phosphate, 15 and a nitrogenous heterocyclic base. The base is linked to the sugar moiety via the glycosidic carbon (1' carbon of the pentose) and that combination of base and sugar is a nucleoside. When the nucleoside contains a phosphate group bonded to the 3' or 5' 20 position of the pentose it is referred to as a nucleotide.

Base Pair (bp): A partnership of adenine (A) with thymine (T), or of cytosine (C) with guanine (G) in a double stranded DNA molecule.

25

B. SP18 Monomer-Containing Compositions

The present invention contemplates a SP18 monomer-containing composition (subject protein composition) wherein the SP18 monomer is present in 30 either substantially isolated or substantially pure form. By "isolated" is meant that SP18 monomer and SP18 dimer are present as part of a composition free of other alveolar surfactant proteins.

By "substantially pure" is meant that SP18 35 monomer is present as part of a composition free of

A DNA segment of the present invention is characterized as consisting essentially of a DNA sequence that encodes a SP18 monomer, preferably human SP18 monomer. That is, a DNA segment of the present invention forms a structural gene capable of expressing a SP18 monomer. While the codons of the DNA segment need not be collinear with the amino acid residue sequence of SP18 monomer because of the presence of an intron, it is preferred that the structural gene be capable of expressing SP18 monomer in mature form, i.e., without the need for post-translational proteolytic processing. Preferably, the gene is present as an uninterrupted linear series of codons where each codon codes for an amino acid residue found in a SP18 monomer, i.e., a gene containing no introns.

Thus, a DNA segment consisting essentially of the sequence shown in Figure 1 from about nucleotide position 187 to about nucleotide position 426, preferably to about nucleotide position 429, and capable of expressing SP18 monomer, constitutes one preferred embodiment of the present invention.

DNA segments that encode SP18 monomer can easily be synthesized by chemical techniques, for example, the phosphotriester method of Matteuccci, et al, J. Am. Chem. Soc., 103:3185 (1981). Of course, by chemically synthesizing the coding sequence, any desired modifications can be made simply by substituting the appropriate bases for those encoding the native amino acid residue sequence.

Also contemplated by the present invention are ribonucleic acid (RNA) equivalents of the above described DNA segments.

sample to be assessed is administered through an endotracheal tube to fetal rabbits or lambs delivered prematurely by Caesarian section. (These "preemies" lack their own PS, and are supported on a ventilator.)

5 Measurements of lung compliance, blood gases and ventilator pressure provide indices of activity. Preliminary assessment of activity may also be made by an in vitro assay, for example that of King, et al, Am. J. Physiol., 223:715-726 (1972), or that

10 illustrated below which utilizes a measurement of surface tension at a air-water interface when a protein or polypeptide is admixed with a phospholipid.

C. Nucleic Acid Segments

15 In living organisms, the amino acid residue sequence of a protein or polypeptide is directly related via the genetic code to the deoxyribonucleic acid (DNA) sequence of the structural gene that codes for the protein. Thus, a structural gene can be

20 defined in terms of the amino acid residue sequence, i.e., protein or polypeptide, for which it codes.

An important and well known feature of the genetic code is its redundancy. That is, for most of the amino acids used to make proteins, more than one

25 coding nucleotide triplet (codon) can code for or designate a particular amino acid residue. Therefore, a number of different nucleotide sequences may code for a particular amino acid residue sequence. Such nucleotide sequences are considered functionally

30 equivalent since they can result in the production of the same amino acid residue sequence in all organisms. Occasionally, a methylated variant of a purine or pyrimidine may be incorporated into a given nucleotide sequence. However, such methylations do not affect

35 the coding relationship in any way.

D. Recombinant Nucleic Acid Molecules

The recombinant nucleic acid molecules of the present invention can be produced by operatively linking a vector to a nucleic acid segment of the 5 present invention.

As used herein, the phase "operatively linked" means that the subject nucleic acid segment is attached to the vector so that expression of the structural gene formed by the segment is under the 10 control of the vector.

As used herein, the term "vector" refers to a nucleic acid molecule capable of replication in a cell and to which another nucleic acid segment can be operatively linked so as to bring about replication of 15 the attached segment. Vectors capable of directing the expression of a structural gene coding for SP18 monomer are referred to herein as "expression vectors." Thus, a recombinant nucleic acid molecule (rDNA or rRNA) is a hybrid molecule comprising at 20 least two nucleotide sequences not normally found together in nature.

The choice of vector to which a nucleic acid segment of the present invention is operatively linked depends directly, as is well known in the art, on the 25 functional properties desired, e.g., protein expression, and the host cell to be transformed, these being limitations inherent in the art of constructing recombinant nucleic acid molecules. However, a vector contemplated by the present invention is at least 30 capable of directing the replication, and preferably also expression, of SP18 monomer structural gene included in a nucleic acid segment to which it is operatively linked.

In preferred embodiments, a vector 35 contemplated by the present invention includes a

(Pharmacia), pBPV-1/pML2d (International Biotechnologies, Inc.), and pTDT1 (ATCC, #31255).

In preferred embodiments, a eucaryotic cell expression vector used to construct an rDNA molecule 5 of the present invention includes a selection marker that is effective in a eucaryotic cell, preferably a drug resistance selection marker. A preferred drug resistance marker is the gene whose expression results in neomycin resistance, i.e., the neomycin 10 phosphotransferase (neo) gene. Southern, et al, J. Mol. Appl. Genet., 1:327-341 (1982).

The use of retroviral expression vectors to form a recombinant nucleic acid molecule of the present invention is also contemplated. As used 15 herein, the term "retroviral expression vector" refers to a nucleic acid molecule that includes a promoter sequence derived from the long terminal repeat (LTR) region of a retrovirus genome.

In preferred embodiments, the expression 20 vector is a retroviral expression vector that is replication-incompetent in eucaryotic cells. The construction and use of retroviral vectors has been described by Sorge, et al, Mol. Cell. Biol., 4:1730-37 (1984).

25 A variety of methods have been developed to operatively link nucleic acid segments to vectors via complementary cohesive termini. For instance, complementary homopolymer tracts can be added to the nucleic acid segment to be inserted and to a terminal 30 portion of the vector nucleic acid. The vector and nucleic acid segment are then joined by hydrogen bonding between the complementary homopolymeric tails to form a recombinant nucleic acid molecule.

35 Synthetic linkers containing one or more restriction sites provide an alternative method of

procaryotic replicon, i.e., a DNA sequence having the ability to direct autonomous replication and maintenance of an rDNA molecule extrachromosomally in a procaryotic host cell, such as a bacterial host cell, transformed therewith. Such replicons are well known in the art. In addition, those embodiments that include a procaryotic replicon also include a gene whose expression confers drug resistance to a bacterial host transformed therewith. Typical bacterial drug resistance genes are those that confer resistance to ampicillin or tetracycline.

Those vectors that include a procaryotic replicon can also include a procaryotic promoter capable of directing the expression (transcription and translation) of a SP18 monomer gene in a bacterial host cell, such as E. coli, transformed therewith. A promoter is an expression control element formed by a DNA sequence that permits binding of RNA polymerase and transcription to occur. Promoter sequences compatible with bacterial hosts are typically provided in plasmid vectors containing convenient restriction sites for insertion of a DNA segment of the present invention. Typical of such vector plasmids are pUC8, pUC9, pBR322 and pBR329 available from Biorad Laboratories, (Richmond, CA) and pPL and pKK223 available from Pharmacia, Piscataway, N.J.

Expression vectors compatible with eucaryotic cells, preferably those compatible with vertebrate cells, can also be used to form an rDNA molecule of the present invention. Eucaryotic cell expression vectors are well known in the art and are available from several commercial sources. Typically, such vectors are provided containing convenient restriction sites for insertion of the desired DNA segment. Typical of such vectors are pSVL and pKSV-10

joining a nucleic acid segment to vectors. For instance, a DNA segment of the present invention is treated with bacteriophage T4 DNA polymerase or E. coli DNA polymerase I, enzymes that remove protruding, 5 3', single-stranded termini with their 3'-5' exonuclease activities and fill in recessed 3' ends with their polymerizing activities. The combination of these activities therefore generates blunt-ended DNA segments. The blunt-ended segments are then 10 incubated with a large molar excess of linker molecules in the presence of an enzyme that is able to catalyze the ligation of blunt-ended DNA molecules, such as bacteriophage T4 DNA ligase. Thus, the products of the reaction are DNA segments carrying 15 polymeric linker sequences at their ends. These DNA segments are then cleaved with the appropriate restriction enzyme and ligated to an expression vector that has been cleaved with an enzyme that produces termini compatible with those of the DNA segment.

20 Synthetic linkers containing a variety of restriction endonuclease sites are commercially available from a number of sources including International Biotechnologies, Inc., New Haven, CT.

Also contemplated by the present invention 25 are RNA equivalents of the above described recombinant DNA molecules.

E. Transformed Cells and Cultures

The present invention also relates to a host 30 cell transformed with a recombinant nucleic acid molecule of the present invention, preferably an rDNA capable of expressing an SP18 monomer. The host cell can be either prokaryotic or eucaryotic.

"Cells" or "transformed host cells" or "host 35 cells" are often used interchangeably as will be clear

from the context. These terms include the immediate subject cell, and, of course, the progeny thereof. It is understood that not all progeny are exactly identical to the parental cell, due to chance 5 mutations or differences in environment. However, such altered progeny are included when the above terms are used.

Bacterial cells are preferred prokaryotic host cells and typically are a strain of E. coli such 10 as, for example, the E. coli strain DH5 available from Bethesda Research Laboratories, Inc., Bethesda, MD. Preferred eukaryotic host cells include yeast and mammalian cells, preferably vertebrate cells such as those from a mouse, rat, monkey or human fibroblastic 15 cell line. Preferred eukaryotic host cells include Chinese hamster ovary (CHO) cells available from the ATCC as CCL61 and NIH Swiss mouse embryo cells NIH/3T3 available from the ATCC as CRL 1658. Transformation of appropriate cell hosts with a recombinant nucleic 20 acid molecule of the present invention is accomplished by well known methods that typically depend on the type of vector used. With regard to transformation of prokaryotic host cells, see, for example, Cohen, et al, Proc. Natl. Acad. Sci. USA, 69:2110 (1972); and 25 Maniatis, et al, Molecular Cloning, A Laboratory Manual, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY (1982). With regard to transformation of vertebrate cells with recombinant nucleic acid molecules containing retroviral vectors, see, for 30 example, Sorge, et al, Mol. Cell. Biol., 4:1730-37 (1984); Graham, et al, Virol., 52:456 (1973); and Wigler, et al, Proc. Natl. Acad. Sci. USA, 76:1373-76 (1979).

Successfully transformed cells, i.e., cells 35 that contain a recombinant nucleic acid molecule of

the present invention, can be identified by well known techniques. For example, cells resulting from the introduction of an rDNA of the present invention can be cloned to produce monoclonal colonies. Cells from 5 those colonies can be harvested, lysed and their DNA content examined for the presence of the rDNA using a method such as that described by Southern, J. Mol. Biol., 98:503 (1975) or Berent, et al, Biotech., 3:208 (1985).

10 In addition to directly assaying for the presence of rDNA, successful transformation can be confirmed by well known immunological methods when the rDNA is capable of directing the expression of an SP18 monomer. For example, cells successfully transformed 15 with an expression vector operatively linked to a DNA segment of the present invention produce proteins displaying SP18 monomer antigenicity. Thus, a sample of a cell culture suspected of containing transformed cells are harvested and assayed for human SP18 using 20 antibodies specific for that antigen, the production and use of such antibodies being well known in the art.

Thus, in addition to the transformed host cells themselves, the present invention also 25 contemplates a culture of those cells, preferably a monoclonal (clonally homogeneous) culture, or a culture derived from a monoclonal culture, in a nutrient medium. Preferably, the culture also contains a protein displaying SP18 monomer 30 antigenicity, and more preferably, biologically active SP18 monomer.

Nutrient media useful for culturing transformed host cells are well known in the art and can be obtained from several commercial sources. In

embodiments wherein the host cell is mammalian, a "serum-free" medium is preferably used.

F. Recombinant Methods for Producing SP18

5 Another aspect of the present invention pertains to a method for producing SP18, preferably human SP18 monomer. The method entails initiating a culture comprising a nutrient medium containing host cells, preferably human cells, transformed with a rDNA 10 molecule of the present invention that is capable of expressing SP18 monomer. The culture is maintained for a time period sufficient for the transformed cells to express SP18 monomer. The expressed protein is then recovered from the culture.

15 Methods for recovering an expressed protein from a culture are well known in the art and include fractionation of the protein-containing portion of the culture using well known biochemical techniques. For instance, the methods of gel filtration, gel 20 chromatography, ultrafiltration, electrophoresis, ion exchange, affinity chromatography and the like, such as are known for protein fractionations, can be used to isolate the expressed proteins found in the culture. In addition, immunochemical methods, such as 25 immunoaffinity, immunoabsorption and the like can be performed using well known methods.

30 Also contemplated by the present invention is an SP18 monomer produced by a recombinant nucleic acid method described herein.

G. Polypeptides

35 A polypeptide of the present invention (subject polypeptide) is characterized by its amino acid residue sequence and novel functional properties. A subject polypeptide when admixed with a

pharmaceutically acceptable phospholipid forms a synthetic pulmonary surfactant having a surfactant activity greater than the surfactant activity of the phospholipid alone (as indicated by a lower ΔP as shown 5 in Figure 6 and a higher volume per given pressure as shown in Figures 7 and 8).

As seen in Figure 1, SP18 has a large hydrophobic region (residues 1 to about 75), followed by a relatively short hydrophilic region at the carboxy 10 terminus (residues 76 to 80, or 81). In referring to amino acid residue numbers of the SP18 sequence, those residues are as illustrated in Figure 1.

In one embodiment, a subject polypeptide consists essentially of at least about 10, preferably 15 at least 11 amino acid residues, and no more than about 60, more usually fewer than about 35 and preferably fewer than about 25 amino acid residues that correspond to the sequence of SP18 monomer.

Usually, the amino acid sequence of a polypeptide 20 of this invention will correspond to a single group of contiguous residues in the linear sequence of SP18. However, polypeptides that correspond to more than one portion of the SP18 sequence are also contemplated. Usually at least one sequence that corresponds to at 25 least 10, preferably at least 15, contiguous residues of the hydrophobic region of SP18 will be present in the peptide. A plurality of hydrophobic region amino acid sequences may be present.

A subject polypeptide will preferably include as 30 its carboxy terminal sequence at least 5 contiguous residues in the linear sequence of SP18 including residue 80. Thus the polypeptides of this invention may include one or more groups of amino acid residues that correspond to portions of SP18 so that a sequence 35 corresponding to a first group of contiguous residues

of the SP18 monomer may be adjacent to a sequence corresponding to a second group of contiguous residues from the same or another portion of the SP18 monomer in the polypeptide sequence. Peptides having two or 5 more sequences that correspond to a single group of contiguous amino acid residues from the linear sequence of SP18 is also contemplated.

Exemplary preferred subject polypeptides corresponding in amino acid residue sequence to human 10 SP18 monomer hydrophobic region are shown in Table 1.

TABLE 1

<u>Designation¹</u>	<u>Amino Acid Residue Sequence</u>
p1-15	FPIPLPYCWLCRALI
p11-25	CRALIKRIQAMIPKG
5 p21-35	MIPKGALAVAVAQVC
p31-45	VAQVCRVVPLVAGGI
p41-55	VAGGICQCLAERYSV
p46-76	CQCLAERYSVILLDTLLGRMLPQLVCRLVLR
p51-65	ERYSVILLDTLLGRM
10 p51-72	ERYSVILLDTLLGRMLPQLVCR
p51-76	ERYSVILLDTLLGRMLPQLVCRLVLR
p54-72	SVILLDTLLGRMLPQLVCR
p54-76	SVILLDTLLGRMLPQLVCRLVLR
p61-75	LLGRMLPQLVCRLVLR

15

1 The designation of each peptide indicates that portion of the amino acid residue sequence of human SP18 monomer, as shown in Figure 1 to which the peptide sequence corresponds, i.e., it

20 indicates the location of the peptide sequence in the protein sequence.

25 In preferred embodiments, a subject polypeptide is further characterized as having a carboxy-terminal amino acid residue sequence represented by the formula:

-RLVLRC_ZSM_DDD₁,

wherein Z is an integer having a value of 0 or 1 such that when Z is 0 the D residue to which it is a

30 subscript is absent and when Z is 1 the D residue to which it is a subscript is present. Exemplary preferred "carboxy-terminal polypeptides" are shown in Table 2.

TABLE 2

<u>Designation</u>	<u>Amino Acid Residue Sequence</u>
p71-81	CRLVLRCSMDD
p66-81	LPQLVCRLVLRCSMDD
5 p59-81	DTLLGRMLPQLVCRLVLRCSMDD
p52-81	RYSVILLDTLLGRMLPQLVCRLVLRCSMDD
P51-81	ERYSVILLDTLLGRMLPQLVCRLVLRCSMDD
P51-80	ERYSVILLDTLLGRMLPQLVCRLVLRCSMDD
p36-81	RVVPLVAGGICQCLAERYSVILLDTLLGRMLPQLVCRLVLRCSMDD
10 p32-81	AQVCRVVPLVAGGICQCLAERYSVILLDTLLGRMLPQLVCRLVLRCSMDD

1 The designation is the same as in Table 1.

15 Preferably, a subject polypeptide has an amino acid residue sequence that corresponds to a portion of the sequence shown in Figure 1. However, it should be understood that a polypeptide of the present invention need not be identical to the amino acid residue sequence of a native SP18 monomer. Therefore, a polypeptide of the present invention can be subject to various changes, such as insertions, deletions and substitutions, either conservative or non-conservative, where such changes provide for 25 certain advantages in their use.

Conservative substitutions are those where one amino acid residue is replaced by another, biologically similar residue. Examples of conservative substitutions include the substitution of 30 one hydrophobic residue such as isoleucine, valine, leucine or methionine for another, or the substitution of one polar residue for another such as between arginine and lysine, between glutamic and aspartic acids or between glutamine and asparagine and the 35 like. The term "conservative substitution" also

includes the use of a substituted amino acid in place of an unsubstituted parent amino acid provided that such a polypeptide also displays the requisite binding activity.

5 In one preferred embodiment, a serine (S) residue is substituted for a cysteine (C) residue, usually at least one of residue positions 71 and 77. Preferably the serine analog has a sequence corresponding to the sequence of residues 51-76 of the 10 SP18 monomer with the substitution at residue 71 or to the sequence of residues 51-81 with serine substitutions at 71 and 77.

When a polypeptide of the present invention has a sequence that is not identical to the sequence 15 of a native SP18 monomer because one or more conservative or non-conservative substitutions have been made, usually no more than about 20 number percent and more usually no more than 10 number percent of the amino acid residues are substituted, 20 except where additional residues have been added at either terminus as for the purpose of providing a "linker" by which the polypeptides of this invention can be conveniently affixed to a label or solid matrix, or carrier. Labels, solid matrices and 25 carriers that can be used with the polypeptides of this invention are described hereinbelow.

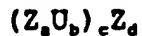
Amino acid residue linkers are usually at least one residue and can be 40 or more residues, more often 1 to 10 residues that do not correspond in amino 30 acid residue sequence to a native SP18 monomer. Typical amino acid residues used for linking are tyrosine, cysteine, lysine, glutamic and aspartic acid, or the like. In addition, a polypeptide sequence of this invention can differ from the natural 35 sequence by the sequence being modified by terminal-

NH, acylation, e.g., acetylation, or thioglycolic acid amidation, terminal-carboxlyamidation, e.g., ammonia, methylamine, etc.

When coupled to a carrier via a linker to
5 form what is known in the art as a carrier-hapten
conjugate, a polypeptide of the present invention is
capable of inducing antibodies that immunoreact with
SP18 monomer. In view of the well established
10 principle of immunologic cross-reactivity, the present
invention therefore contemplates antigenically related
variants of the polypeptides shown in Tables 1 and 2.
An "antigenically related variant" is a polypeptide
that includes at least a six amino acid residue
sequence portion of a polypeptide from Table 1 or
15 Table 2 and which is capable of inducing antibody
molecules that immunoreact with a polypeptide from
Table 1 or 2 and an SP18 monomer.

In another embodiment, a polypeptide of this
invention has amino acid residue sequence that has a
20 composite hydrophobicity of less than zero, preferably
less than or equal to -1, more preferably less than or
equal to -2. Determination of the composite
hydrophobicity value for a peptide is described in
detail in Example 2. These hydrophobic polypeptides
25 perform the function of the hydrophobic region of
SP18. In a preferred embodiment, the amino acid
sequence mimics the pattern of hydrophobic and
hydrophilic residues of SP18.

A preferred hydrophobic polypeptide includes a
30 sequence having alternating hydrophobic and
hydrophilic amino acid residue regions and is
characterized as having at least 10 amino acid
residues represented by the formula:



Z and U are amino acid residues such that at each occurrence Z and U are independently selected. Z is a hydrophilic amino acid residue, preferably selected from the group consisting of R, D, E and K. U is a hydrophobic amino acid residue, preferably selected from the group consisting of V, I, L, C, Y and F. "a", "b", "c" and "d" are numbers which indicate the number of hydrophilic or hydrophobic residues. "a" has an average value of about 1 to about 5, preferably about 1 to about 3. "b" has an average value of about 3 to about 20, preferably about 3 to about 12, most preferably, about 3 to about 10. "c" is 1 to 10, preferably 2 to 10, most preferably 3 to 6. "d" is 1 to 3, preferably 1 to 2.

By stating that the amino acid residue represented by Z and U is independently selected, it is meant that at each occurrence a residue from the specified group is selected. That is, when "a" is 2, for example, each of the hydrophilic residues represented by Z will be independently selected and thus can include RR, RD, RE, RK, DR, DD, DE, DK, etc. By stating that "a" and "b" have average values, it is meant that although the number of residues within the repeating sequence (Z, U_b) can vary somewhat within the peptide sequence, the average values of "a" and "b" would be about 1 to about 5 and about 3 to about 20, respectively.

Exemplary preferred polypeptides of the above formula are shown in Table 3.

TABLE 3

<u>Designation¹</u>	<u>Amino Acid Residue Sequence</u>
DL4	DLLLLDLLLLDLLLLDLLLLD
RL4	RLLLLRLLLLRLLLLRLLLLR
5 RL8	RLLLLLRLRLRLRLRLRL
RL7	RRLLLLLRLRLRLRLRLRL
RCL1	RLLLLCLLRLRLRLRL
RCL2	RLLLLCLLRLRLRLRL
RCL3	RLLLLCLLRLRLRLRLRL

10

1 The designation is an abbreviation for the indicated amino acid residue sequence.

Also contemplated are composite polypeptides of 10 to 60 amino acid residues. A composite polypeptide consists essentially of an amino terminal sequence and a carboxy terminal sequence. The amino terminal sequence has an amino acid sequence of a hydrophobic region polypeptide or a hydrophobic peptide of this invention, preferably hydrophobic polypeptide, as defined in the above formula. The carboxy terminal sequence has the amino acid residue sequence of a subject carboxy terminal peptide.

A polypeptide of the present invention can be synthesized by any techniques that are known to those skilled in the polypeptide art. An excellent summary of the many techniques available may be found in J.M. Steward and J.D. Young, "Solid Phase Peptide Synthesis", W.H. Freeman Co., San Francisco, 1969, and J. Meienhofer, "Hormonal Proteins and Peptides", Vol. 2, p. 46, Academic Press (New York), 1983 for solid phase peptide synthesis, and E. Schroder and K. Kubke, "The Peptides", Vol. 1, Academic Press (New York), 1965 for classical solution synthesis.

35 In general, these methods comprise the sequential addition of one or more amino acid residues

or suitably protected amino acid residues to a growing peptide chain. Normally, either the amino or carboxyl group of the first amino acid residue is protected by a suitable, selectively removable protecting group. A 5 different, selectively removable protecting group is utilized for amino acids containing a reactive side group such as lysine.

Using a solid phase synthesis as exemplary, the protected or derivatized amino acid is attached to 10 an inert solid support through its unprotected carboxyl or amino group. The protecting group of the amino or carboxyl group is then selectively removed and the next amino acid in the sequence having the complimentary (amino or carboxyl) group suitably 15 protected is admixed and reacted under conditions suitable for forming the amide linkage with the residue already attached to the solid support. The protecting group of the amino or carboxyl group is then removed from this newly added amino acid residue, 20 and the next amino acid (suitably protected) is then added, and so forth. After all the desired amino acids have been linked in the proper sequence, any remaining terminal and side group protecting groups (and solid support) are removed sequentially or 25 concurrently, to afford the final polypeptide.

H. Synthetic Surfactants

Recombinantly produced SP18 and/or a subject polypeptide can be admixed with a pharmaceutically 30 acceptable phospholipid to form a synthetic pulmonary surfactant (PS) useful in the treatment of respiratory distress syndrome.

The phase "pharmaceutically acceptable" refers to molecular entities and compositions that do

not produce an allergic or similar untoward reaction when administered to a human.

5 Phospholipids useful in forming synthetic alveolar surfactants by admixture with protein are well known in the art. See, Notter, et al, Clin. Perinatology, 14:433-79 (1987), for a review of the use of both native and synthetic phospholipids for synthetic surfactants.

In one embodiment, the present invention 10 contemplates a synthetic pulmonary surfactant effective in treating RDS comprising an effective amount of a subject polypeptide admixed with a pharmaceutically acceptable phospholipid. While methods for determining the optimal 15 polypeptide:phospholipid weight ratios for a given polypeptide-phospholipid combination are well known, therapeutically effective ratios are in the range of about 1:5 to about 1:10,000, preferably about 1:100 to about 1:5,000, and more preferably about 1:500 to 20 about 1:1000. In a more preferred embodiment, the polypeptide:phospholipid weight ratio is in the range of about 1:5 to about 1:2,000, preferably about 1:7 to about 1:1,000, and more preferably about 1:10 to about 1:100. Thus, a synthetic pulmonary surfactant of this 25 invention can contain about 50, usually about 80, to almost 100 weight percent lipid and about 50, usually about 20, to less than 1 weight percent polypeptide. Preferably a subject polypeptide is about 1 to about 10 weight percent of the surfactant for polypeptides 30 corresponding to portions of the SP18 sequence and 1:100 for polypeptides corresponding to the entire SP18 monomer.

The lipid portion is preferably about 50 to about 90, more preferably about 50 to about 75, weight 35 percent dipalmitoylphosphatidylcholine (DPPC) with the

remainder unsaturated phosphatidyl choline, phosphatidyl glycerol (PG), triacylglycerols, palmitic acid sphingomyelin or admixtures thereof. The synthetic pulmonary surfactant is prepared by admixing

5 a solution of a subject polypeptide with a suspension of liposomes or by admixing the subject polypeptide and lipids directly in the presence of organic solvent. The solvent is then removed by dialysis or evaporation under nitrogen and/or exposure to vacuum.

10 A subject synthetic pulmonary surfactant is preferably formulated for endotracheal administration, e.g., typically as a liquid suspension, as a dry powder "dust", or as an aerosol. For instance, a synthetic surfactant (polypeptide-lipid micelle) is

15 suspended in a liquid with a pharmaceutically acceptable excipient such as water, saline, dextrose, glycerol and the like. A surfactant-containing therapeutic composition can also contain small amounts of non-toxic auxiliary substances such as pH buffering

20 agents including sodium acetate, sodium phosphate and the like. To prepare a synthetic surfactant in dust form, a synthetic surfactant is prepared as described herein, then lyophilized and recovered as a dry powder.

25 If it is to be used in aerosol administration, a subject synthetic surfactant is supplied in finely divided form along with an additional surfactant and propellant. Typical surfactants which may be administered are fatty acids

30 and esters. However, it is preferred, in the present case, to utilize the other components of the surfactant complex DPPC and PG. Useful propellants are typically gases at ambient conditions, and are condensed under pressure. Lower alkane and

35 fluorinated alkane, such as Freon, may be used. The

aerosol is packaged in a container equipped with a suitable valve so that the ingredients may be maintained under pressure until released.

A synthetic surfactant is administered, as appropriate to the dosage form, by endotracheal tube, by aerosol administration, or by nebulization of the suspension or dust into the inspired gas. Amounts of synthetic PS between about 1.0 and about 400 mg/kg, preferably about 50 mg to about 500 mg/kg, are administered in one dose. For use in newly born infants, one to three administrations are generally sufficient. For adults, sufficient reconstituted complex is administered to produce a PO, within the normal range (Hallman, et al, J. Clinical Investigation, 70:673-682, 1982).

The following examples are intended to illustrate, but not limit, the present invention.

20

Examples

Example 1 - Isolation and Characterization of Native SP18

A. Methods

Purification of LMW apoproteins

25 Human pulmonary surfactant was isolated from full-term amniotic fluid and applied to a column of DEAE-Sephacel A-50 (Pharmacia, Uppsala, Sweden) using 4 milliliter (ml) packed volume per 200 milligram (mg) surfactant, in a tris-EDTA buffer containing 1% n-octyl-beta-D-glucopyranoside as described by Revak, et al, Am. Rev. Respir. Dis., 134:1258-1265 (1986) and Hallman, et al, Pediatrics, 71:473-482 (1983). This particular column and conditions were used in order to isolate the 35,000 dalton apoprotein (for use in other 30 studies) without exposing it to potentially denaturing

organic solvents. The void volume, containing the lipids and proteins which did not bind to the column under these conditions, was pooled and extracted with an equal volume of 2:1 chloroform:methanol.

5 Following centrifugation to separate the phases, the upper phase (water + methanol) was re-extracted with 1/2 volume chloroform. After centrifugation, the resultant lower organic phase was added to the initial lower phase and evaporated to dryness under a stream 10 of nitrogen. This extract, which contained 100-180 mg phospholipid, LMW apoproteins, and octylglucopyranoside, was redissolved in 2.5 ml of chloroform:methanol, 2:1.

Following the method of Takahashi, et al, Biochem. 15 Biophys. Res. Comm., 135:527-532 (1986), which was found to afford a good separation of octylglucopyranoside from the LMW proteins and phospholipids, a glass column 2.5 cm in diameter was packed at 4° to a height of 38 cm with Sephadex LH-20 20 (Pharmacia, Uppsala, Sweden) in 2:1 chloroform:methanol. The sample was loaded and 2 ml fractions collected as chloroform:methanol, 2:1, was run through at a flow rate of 8.5 ml/hr. Phospholipids eluted after 40 ml of buffer had passed through the 25 column. Octylglycopyranoside appeared at the 56-116 ml region.

The phospholipid region was pooled, dried under nitrogen, and redissolved in 1 ml chloroform. A silicic acid column was prepared by packing 9 ml of 30 Bio-Sil HA (BioRad, Richmond, CA) in chloroform in a glass column at room temperature. The sample (which contained approximately 50 mg phospholipid) was applied and washed with 11 ml chloroform. A linear gradient of increasing methanol was established using 35 an equal weight of chloroform and methanol (38.8 g,

26.5 ml chloroform and 50 ml methanol). Fractions of 2 ml were collected as the gradient was applied to the column. Figure 1 shows the protein and phospholipid profiles obtained.

5 Phospholipid analyses showed a small peak in fractions 17-20 and a major peak after fraction 30. The Pierce BCA protein assay was positive in fractions 12-19 and 28-33, but it should be noted that the latter peak is likely to be due to the phospholipid

10 15 present in this region. Electrophoresis in sodium dodecyl sulfate polyacrylamide gels showed the LMW apoproteins were present in fractions 13-19 with some separation occurring between SP9 and SP18.

Alternatively, a method devised by Hawgood, et al, 15 Proc. Natl. Acad. Sci., 85:66-70 (1987) employing a butanol extraction of PS followed by chromatography of Sephadex LH-20 in an acidified chloroform/methanol buffer, can be used to isolate the LMW apoprotein mixture. For some studies, a separation of the two 20 LMW apoproteins was effected using Sephadex LH-60. A glass column of 1 cm diameter was packed to 40 cm with Sephadex LH-60 (Pharmacia, Uppsala, Sweden) in chloroform/methanol, 1:1, containing 5% 0.1 N HCl. A flow rate of 1-2 ml/hr was used. A mixture of the LMW 25 apoproteins containing about 200-700 micrograms (ug) of protein from either the Bio-Sil HA column or the LH-20 column described by Hawgood, et al, Proc. Natl. Acad. Sci., 85:66-70 (1987), in a volume of 0.5 ml buffer, was applied to the top of the column and 30 fractions of 0.5 ml were collected. Typically, SP18 protein eluted in fractions 16-19 and SP9 in fractions 24-29. Appropriate fractions were pooled and dried in glass tubes under nitrogen. A brief period of lyophilization ensured complete removal of the HCl. 35 Proteins were re-solubilized in methanol prior to use.

SDS-Gel Electrophoresis

Gel electrophoresis in 16% polyacrylamide was performed in the presence of sodium dodecyl sulfate 5 (SDS-PAGE) according to the method of Laemmli, Nature, 227:680-685 (1970), using 3x7 cm minislab gels. 1% β -mercaptoethanol was added to samples where indicated as a disulfide reducing agent. Following 10 electrophoresis, the gels were fixed overnight in 50% methanol + 12% acetic acid, washed in water for 2 hours, and silver-stained according to the method of Wray, et al, Anal. Biochem., 118:197-203 (1981).

Octylglucopyranoside Assay

15 An assay for the quantitation of n-octyl-beta-D-glyco-pyranoside, based on the anthrone method of Spiro, Methods Enzymol., 8:3-5 (1966) has been described previously by Revak, et al, Am. Rev. Respir. Dis., 134:1258-1265 (1986).

20

Protein Determinations

Organic samples containing up to 5 μ g protein were dried in 12x75 mm glass tubes under nitrogen. Fifteen microliters (ul) of 1% SDS in H_2O and 300 ul BCA 25 Protein Assay Reagent (Pierce Chemical Co., Rockford, IL) were admixed with the protein in each tube. Tubes were covered and incubated at 60°C for 30 min. After cooling, the samples were transferred to a 96-well flat-bottom polystyrene microtiter plate and OD₅₅₀ 30 measured. Bovine serum albumin was used as a standard. It should be noted that some phospholipids will react in the BCA protein assay making protein quantitations inaccurate when lipid is present (i.e., prior to Bio-Sil HA chromatography). Additionally, 35 once purified, the hydrophobic LMW apoproteins

themselves react poorly with the BCA reagents and all quantitations of the isolated proteins were, therefore, based on amino acid compositions.

5 Phospholipids

Dipalmitoylphosphatidylcholine (DPPC, beta, gamma-dipalmitoyl-L-alpha-lecithin) and L-alpha-phosphatidyl-DL-glycerol (PG, derivative of egg lecithin) were purchased from either Calbiochem-
10 Behring (La Jolla, CA) or Avanti Polar-Lipids, Inc. (Birmingham, AL). DPPC was added to PG in chloroform in a weight ratio of 3:1.

Admixture of LMW Apoproteins with Phospholipids

15 For in vitro assays, a methanol solution containing 4 ug of SP9 or SP18, was added to 400 ug DPPC:PG in chloroform in a 12x75 mm glass tube. Following a brief vortex mixing, the samples were dried under N₂. Ninety microliters of water were added
20 to each and the tubes placed in a 37°C water bath for 15 minutes, with periodic gentle mixing. Isotonicity was restored with the addition of 10 ul of 9% NaCl to each sample prior to assay. For in vivo rabbit studies, 50 ug LMW apoproteins (containing both SP9
25 and SP18) or 25 ug SP9 or 25 ug SP18 were dried under N₂. Five mg of phospholipids (DPPC:PG, 3:1) were added in chloroform. The samples were mixed, dried, and resuspended in 250 ul 100 millimolar (mM) saline containing 1.5 mM CaCl₂, to yield a reconstituted
30 surfactant at 20 mg/ml with 0.5-1% protein.

Surfactant Activity Assays

In vitro assays of surfactant activity, assessed as its ability to lower the surface tension of a
35 pulsating bubble, and in vivo assays utilizing fetal

rabbits, have both been described in detail previously by Revak, et al, Am. Rev. Respir. Dis., 134:1258-1265 (1986).

5 Morphometric Analyses

Fetal rabbit lungs, inflated to 30 cm H₂O and then deflated to 10 cm H₂O, were submerged in 10% formalin for 72 hours. Paraffin sections were oriented from apex to base and 5 micron sections taken anterior to 10 posterior. After hematoxylin and eosin staining, 10 fields (100 x) were point-counted from apex to base on multiple sections. Standardized morphometric methods (Weiber, in "Stereological Methods," Vol. I, Academic Press, New York, pp. 33-58, 1979) were used to 15 determine ratios of lung interstitium to air spaces for each treatment group. Intersections of alveolar perimeters were also determined.

Phospholipid Phosphorus Assays

20 Phospholipids were quantitated according to the method of Bartlett, J. Biol. Chem., 234:466-468 (1959).

Amino Acid Analysis

25 Triplicate samples for amino acid compositions were hydrolyzed with HCl at 110°C for 24 hours, with HCl at 150°C for 24 hours, or in performic acid at 110°C for 24 hours followed by HCl hydrolysis at 110°C for 24 hours. Analyses were performed on a Beckman 30 model 121-M amino acid analyzer (Beckman Instruments, Fullerton, CA). Tryptophan was not determined.

Amino Acid Sequencing

35 Vapor-phase protein sequencing was performed on an Applied Biosystems 470A Amino Acid Sequencer (Applied

Biosystems, Inc., Foster City, CA) with an on-line Model 120A HPLC.

Isolation of cDNA Clones for Human SP18

5 RNA was prepared according to Chirgwin, et al, Biochemistry, 18:5294-5299 (1979) from a sample of unaffected adult lung tissue obtained during surgical removal of a neoplastic lesion. Preparation of double stranded cDNA was carried out using standard
10 techniques (Chirgwin et al., supra, and Efstratiadis et al., in "Genetic Engineering", eds. Stelow and Hollaender, Plenum, New York, 1:15-49 (1979) and a library was constructed in lambda NM607 as described by Le Bouc, et al, F.E.B.S. Letts., 196:108-112
15 (1986). SP18 clones were identified by screening phage plaques with synthetic oligonucleotide probes (Benton, et al, Science, 196:180-182 (1977) which were prepared using an Applied Biosystems automated synthesizer and purified by HPLC. Initial candidate
20 clones were obtained using probe TG996 (5' CATTGCCTGTGGTATGCCCTGCCTCC 3') which was derived from the partial nucleotide sequence of a small human surfactant apoprotein cDNA (Schilling et al., International Patent Application WO 86/03408). Larger
25 clones (up to 1.5 kb) were isolated using probe TG1103 (5' TCGAGCAGGATGACGGAGTAGCGCC 3') which was based on the 5' sequence of one of the original clones. The nucleotide sequence of the cDNA clones was determined by the chain termination method (Sanger, et al, Proc. Natl. Acad. Sci. U.S.A., 74:5463-5467 (1977) using EcoRI restriction fragments subcloned in an appropriate M13 vector.

B. Results

Characteristics of the LMW Apoproteins

The LMW apoproteins isolated from human amniotic fluid appeared after silicic acid chromatography, or 5 after the Sephadex LH-20 column chromatography Figure 2 described by Hawgood, et al, Proc. Natl. Acad. Sci. 85:66-70 (1987), as two protein bands in SDS- polyacrylamide gel electrophoresis under non-reducing conditions. The upper band, having an apparent 10 molecular weight of 18,000 daltons is a dimer, and therefore designated SP18 dimer. With the addition of β -mercaptoethanol, SP18 dimer reduced to 9,000 daltons and was designated SP18 monomer (Fig. 3). The other LMW apoprotein, designated SP9, appears as a diffuse 15 band between 9 and 12,000 daltons in the presence or absence of reducing agents. SP9 was separated from SP18 dimer and SP18 monomer by chromatography on Sephadex LH-60. The resultant purified proteins are shown in Fig. 3.

20 Amino acid compositions were determined for SP18 monomer and SP9. Because of the extremely hydrophobic nature of these proteins, HCl hydrolysis was performed at 150°C for 24 hours, in addition to the standard 110°C for 24 hour hydrolysis, and values for valine, 25 leucine, and isoleucine calculated from analyses of the hydrolysates done under the extreme conditions. As shown in Table 3, both proteins are extremely hydrophobic with high levels of valine and leucine.

TABLE 3

5 Amino Acid Composition of Human SP9 and SP18 monomer
and a Comparison with the Theoretical Composition of
SP18¹

	<u>Amino Acid</u>	<u>SP9 (mole %)</u>	<u>SP18 (mole %)</u>	<u>SP18² (mole %)</u>
10	Aspartic acid (or Asparagine)	1.1	3.4	3.7
	Threonine	0.8	1.5	1.2
15	Serine	1.8	2.7	2.5
	Glutamic acid (or Glutamine)	1.5	6.7	6.2
20	Proline	8.3	7.8	7.4
	Glycine	10.6	6.1	4.9
25	Alanine	4.9	10.2	9.9
	Cysteine ²	9.1	7.2	8.6
	Valine ³	12.2	11.7	11.1
30	Methionine	3.4	3.2	3.7
	Isoleucine ³	6.8	6.4	7.5
35	Leucine ³	22.4	17.4	17.3
	Tyrosine	0.7	2.2	2.5
	Phenylalanine	2.6	1.5	1.2
40	Histidine	5.4	0.0	0.0
	Lysine	4.7	3.0	2.5
45	Arginine	3.9	9.0	8.6
	Tryptophan	N.D. ⁴	N.D. ⁴	1.2

50 1 The theoretical composition is based on
sequence data through residue 81.

2 Cysteine content was determined following
performic acid and HCl hydrolyses.

3 Isoleucine and leucine content were each determined following 24 hour HCl hydrolysis at 150°C.

5 4 Tryptophan was not determined.

Amino-terminal sequence analysis of SP18 monomer yielded the following sequence:

10 $\text{NH}_2\text{-Phe-Pro-Ile-Pro-Leu-Pro-Try-}$.

Repeated sequencing of the purified SP9 monomer showed multiple peptides, all rich in leucine and containing at least six consecutive valines. NH_2 -terminal analysis showed phenylalanine, glycine, and 15 isoleucine with the relative amounts of each varying from preparation to preparation.

Nucleotide Sequence Analysis of SP18 cDNA

The nucleotide sequence of a SP18 monomer cDNA 20 clone is presented in Figure 1. The sequence displays 83% homology with the canine SP18 cDNA (Hawgood, et al, Proc. Natl. Acad. Sci., 85:66-70 (1987)). A sequence within a large open reading frame was identified which matches perfectly with the amino 25 terminus of SP18 monomer as determined by Edman degradation of the isolated protein (underlined in Figure 3). This suggests that mature SP18 monomer arises by processing of a larger precursor molecule. In the mature sequence there is a single potential N- 30 glycosylation site (Asn 110), no sites for tyrosine sulfation, and no G-X-Y repeats as found in the 35,000 dalton apoprotein (White, et al, A. Pediatrics Research, 19:501-508; 1985).

The molecular weight of 9000 daltons obtained by 35 SDS-PAGE of reduced SP18 dimer is lower than that predicted for the complete precursor protein sequence with amino terminus $\text{NH}_2\text{-Phe-Pro-Ile-Pro-Leu-Pro-Tyr}$

(19,772 daltons), implying further processing in the region of amino acids 70-90. In support of this, the theoretical amino acid composition (Table I) of a putative 9000 dalton protein comprising residues 1 to 5 81 compares well with the determined values for purified SP18 monomer. The amino terminal portion of the precursor protein (residues 1 to 81) is alkaline and more hydrophobic than the COOH terminal portion (residues 82 to 181): the Kyte-Doolittle index for 10 residues 1 to 81 is 9100 (pI 8.6) and is -3000 (pI 5.91) for residues 82 to 181 (Kyte, et al, J. Pediatrics, 100:619-622; 1982). The amino terminus (residues 1 to 81) is, as in the canine sequence (Hawgood, et al, Proc. Natl. Acad. Sci., 85:66-70; 15 1987), composed of three hydrophobic domains: residues 1 to 11, 22 to 49 and 53 to 74. These are interspersed with a charged domain (residues 12 to 21) and two hydrophilic and charged stretches (residues 47 to 54 and 72 to 81).

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Reconstitution of Surfactant Activity with LMW Apoproteins

Samples were prepared containing 400 ug/100 ul phospholipids (DPPC:PG, 3:1 by weight), phospholipids 25 plus 4 ug SP9, or phospholipids plus 4 ug SP18. Each sample was assayed in the pulsating bubble surfactometer for the ability to lower surface tension. The results are shown in Table 4 as the mean minimal surface tension at 15 sec, 1 minute (min), and 30 5 min. Natural human surfactant, isolated from term amniotic fluid, diluted to 4 mg/ml is shown for comparison. While neither phospholipids nor LMW apoproteins alone had significant surface-tension lowering capacities, a mixture of phospholipids with 35 either SP9 or SP18 showed significant activity.

Recombining the phospholipids with 1% by weight of SP18 lowered the surface tensions measured to levels comparable to those obtained with an equal amount of natural human surfactant (6.3 ± 0.2 dynes/cm for 5 phospholipids + SP18 at 15 sec, 2.0 ± 1.2 dynes/cm for natural surfactant). On an equal weight basis, SP9 lowered surface tension less effectively (16.7 ± 0.8 dynes/cm at 15 sec).

10

TABLE 4Minimum Surface Tensions in the Pulsating Bubble¹

	15 sec	1 min	5 min
PL ²	42.9 ± 1.4	41.6 ± 1.6	34.9 ± 4.9
PL + SP9 ³	16.7 ± 0.8	14.1 ± 1.2	12.2 ± 1.0
PL + SP18 ³	6.3 ± 0.2	5.1 ± 1.0	4.9 ± 0.6
natural human surfactant ⁴	2.0 ± 1.2	2.4 ± 1.4	0.4 ± 0.4

1 Pulsation of 20 cycles/min started 10 sec 20 after bubble formation. All values are in dynes·cm⁻¹ and are the average of at least 3 determinations.

2 Phospholipids DPPC:PG, 3:1, 4 mg/ml

3 1% by weight compared to phospholipids

4 diluted to 4 mg/ml

30

In vivo assays of exogenous (synthetic) surfactant activity were performed by instilling into the airways of immature fetal rabbits saline solutions containing Ca²⁺ alone or with the addition of 35 phospholipids, phospholipids plus LMW apoproteins, or natural human surfactant. The animals were ventilated for 30 min and then degassed by placement in a bell jar under vacuum. The lungs were then inflated to given pressures and the volume of air required for

each pressure was noted. The volumes required for given pressures during deflation from 30 cm H₂O were likewise determined. The resulting pressure/volume curves are shown in Fig. 4 for animals which received 5 synthetic surfactant made with purified SP9 or SP18 (0.5% by weight compared with total phospholipid concentration) and appropriate control animals. Improved lung compliance is apparent in those animals treated with natural or either synthetic surfactant as 10 compared with those receiving saline or phospholipids with the SP18 appearing more effective than SP9 on an equal weight basis. A similar study was performed using a mixture of SP9 and SP18. The results were almost identical to the phospholipid plus SP18 curve 15 presented in Fig. 4.

Following compliance measurements, the lungs were inflated to 30 cm H₂O, deflated back to 10 cm H₂O, clamped, excised and fixed in formalin. Thin sections 20 were stained with hematoxylin and eosin and examined microscopically. As shown in Fig. 5, lungs treated with saline (A) or phospholipids (C) appeared atelectatic while those from animals which received natural (B) or reconstituted (D) surfactant showed normal alveolar expansion. Morphometric analyses of 25 the thin sections showed an interstitium to air space ratio of 4.70 for saline treatment and 3.29 for phospholipids alone as compared with 0.498 for natural surfactant and 0.538 for reconstituted surfactant. These data are shown in Table 5 and corroborate the 30 significant ($p<0.001$; Mann-Whitney U Test) increase in air space seen in Fig. 5. A comparison of alveolar perimeters similarly demonstrated a significantly ($p<0.003$) greater number of intersections of the 35 alveolar boundaries in saline- or phospholipid-treated fetuses compared to surfactant-treated animals.

TABLE 5
 Morphometric Analysis of Airspace
 Following Fetal Rabbit Treatment

	<u>Tracheal Instillation</u>	<u>Interstitium/Air Space</u>
5	saline	4.70
10	phospholipids ¹	3.29
15	phospholipids ¹ + LMW Apoproteins ²	0.538
20	natural human surfactant ³	0.498
	1 2 mg of 3:1 DPPC:PG per animal	
	2 20 ug of LMW apoproteins added to phospholipids	
25	3 2 mg per animal	

C. Discussion

25 This study describes two low molecular weight apoproteins isolated from human amniotic fluid surfactant which can be added to known phospholipids to produce a biologically active pulmonary surfactant. While the proteins in the current study have been 30 designated as SP18 dimer, SP18 monomer and SP9, it is apparent from the recent literature that multiple nomenclature and an assortment of reported molecular weights for LMW PS apoproteins (ranging from 5-18,000 daltons) exist. The apparent differences in physical 35 properties may be explained by a variety of factors including species differences, varying purification and handling techniques, varying determinations of low molecular weights based on standards in SDS-polyacrylamide gels, and potential interference by 40 lipids of low molecular weight protein bands in gels.

Comparisons of amino acid compositions and sequences and immunologic analyses using monospecific antibodies will help to sort out the LMW apoproteins.

It is felt that the SP9 protein described 5 herein, giving a diffuse band on SDS-polyacrylamide gels from 9-12,000 daltons under reducing or non-reducing conditions, is probably the same protein as that designated SAP-6 by Whitsett, et al, Pediatric Research, 20:744-749 (1986), SP5-8 by Hawgood, et al, 10 Proc. Natl. Acad. Sci., 85:66-70 (1987), PSP-6 by Phelps, et al, Am. Rev. Respir. Dis., 135:1112-1117 (1987), and the 5 kDa proteolipid of Takahashi, et al, Biochem. Biophys. Res. Comm., 135:527-532 (1986). The extremely hydrophobic nature of this protein is 15 apparent from its amino acid composition (Table 3) and sequence data, showing at least six consecutive valine residues preceded by a leucine-rich region. The presence of three amino-terminal residues (phenylalanine, glycine, and isoleucine) in the 20 preparations of SP9 derived herein from amniotic fluid surfactant suggests a collection of peptides having an identical sequence but having had one or two residues removed from the amino-terminus. Phelps, et al, Am. Rev. Respir. Dis., 135:1112-1117 (1987) have recently 25 reported a similar finding with bovine PSP-6 apoprotein.

SP18 dimer is comprised of two identical 9000 dalton peptides (but different from the 9000 dalton peptide of SP9) that are disulfide linked. The amino 30 acid composition of SP18 monomer (Table 3) shows a high number of hydrophobic residues. When unreduced SDS-PAGE were overloaded with SP18 protein, sequentially less intensely staining bands were seen at 36,000 and 56,000 daltons, suggesting oligomeric

forms of the protein; upon reduction, only a single 9000 dalton band was seen.

Both SP9 and SP18 dimer apoproteins isolated as described above, could be shown to have biophysical 5 activity following recombination with phospholipids. The addition of 1% by weight of SP18 dimer to the phospholipids DPPC:PG resulted in an immediate increase in surface pressure resulting in surface 10 tensions of less than 10 dynes/cm by 15 sec. The addition of 1% SP9 to DPPC:PG was slightly less 15 effective, lowering surface tensions to 16.7, 14.1, and 12.2 dynes/cm at 15 sec, 1 and 5 min, respectively. Mixtures of both SP18 dimer and SP9 were also effective but further studies will be required to determine whether the combined effect is additive or synergistic.

In vivo studies of reconstituted surfactant using the fetal rabbit model (Schneider, et al, J. Pediatrics, 100:619-622; 1982) were performed using 20 mixtures of SP18 dimer and SP9 as well as each protein individually. A marked improvement in lung compliance was seen in animals treated with natural surfactant or reconstituted surfactant prepared with SP18 dimer apoprotein, as compared with those receiving 25 phospholipids alone or saline (Fig. 4). A moderate improvement was seen when SP9 was used. Identical studies using a mixture of SP18 dimer and SP9 to prepare the reconstituted surfactant showed results very similar to those obtained with SP18 dimer alone 30 (solid squares, Fig. 4); however, the exact ratio of SP18 dimer and SP9 in those studies could not be accurately ascertained. Fig. 5 shows representative microscopic alveolar fields indicating the lack of atelectasis following surfactant instillation.

Suzuki, et al, (Eur. J. Respir. Dis., 69:336-345; 1986) have reported a reduction in surface tension (measured on the Wilhelmy balance or in a pulsating bubble) and a five fold increase in tidal volumes of prematurely-delivered rabbits at insufflation pressures of 25 cm and H₂O when porcine LMW (<15,000 daltons) surfactant apoproteins are added to mixtures of DPPC:PG) at a weight ratio of 5:80:20 (protein:DPPC:PG). Whether one or multiple proteins are present in this system is unclear.

Previous studies using the 35,000 dalton apoprotein (Revak, et al, Am. Rev. Respir. Dis., 134:1258-1265; 1986) also showed moderate reduction in surface tension, similar to that obtained with SP9 in the studies described herein. Clearly, further studies must be done using various combinations and concentrations of SP18, SP9 and the 35,000 dalton apoprotein, as well as Ca⁺⁺ and perhaps various phospholipids to elucidate the interactions between these various components of surfactant and to determine the best conditions for a biologically active exogenous surfactant.

Example 2 - In Vitro Assessment of Polypeptide

Surfactant Activity

A. Methods

Measurement of Surfactant Activity

Measurements of surface pressure across an air-liquid interface (expressed in negative cm of H₂O pressure) at minimal bubble radius were determined at various times using the pulsating bubble technique described by Enhorn, J. Appl. Physiol., 43:198-203 (1977).

Briefly, the Enhorn Surfactometer

(Surfactometer International, Toronto, Ontario)

measures the pressure gradient (ΔP) across a liquid-air interface of a bubble that pulsates at a rate of 20 cycles/min between a maximal (0.55 mm) and minimal (0.4 mm) radius. The bubble, formed in a 37°C, water-
5 enclosed, 20-ul sample chamber, is monitored through a microscopic optic while the pressure changes are recorded or a strip chart recorder calibrated for 0 and -2 cm H₂O.

10 Determination of Composite Hydrophobicity Value

The composite hydrophobicity value of each peptide was determined by assigning to each amino acid residue in a peptide its corresponding hydrophilicity value as described in Table 1 of Hopp, et al, Proc. 15 Natl. Acad. Sci., U.S.A., 78:3824-3829 (1981), which disclosure is incorporated herein by reference. For a given peptide, the hydrophilicity values were summed, the sum representing the composite hydrophobicity value.

20

Preparation of Synthetic Surfactants

After admixture with solvent, each peptide was combined with phospholipids (DPPC:PG), 3:1 to form a synthetic surfactant according to one of the following 25 methods.

Method A was accomplished by admixing 16 ul of peptide/solvent admixture (40 ug peptide) with 100 ul of chloroform containing 400 ug phospholipids, agitating the admixture for about 10 min. at 37°C to 30 form a peptide/phospholipid admixture. Chloroform was removed from the peptide/phospholipid admixture by drying under N₂. The synthetic surfactant thus formed was then admixed with 90 ul of H₂O and gently agitated for about 10 minutes at 37°C. Subsequently, 10 ul of

9% NaCl was admixed to the surfactant containing solution.

Method B was accomplished by first placing 100 ul of chloroform containing 400 ug of phospholipids in a

5 glass tube and removing the chloroform by drying under N₂ for about 10 minutes at 37°C. Sixteen ul of peptide/solvent admixture and 74 ul H₂O were admixed with the dried phospholipids, and then gently agitated for about 10 minutes at 37°C. To the synthetic

10 surfactant thus formed was admixed 10 ul of 9% NaCl.

Method C was accomplished by first maintaining the polypeptide-PL admixture at 43°C for 10 minutes, after which time the solvents were removed from the admixture by drying under N₂. When needed, admixtures

15 were further dried by 15 minutes exposure to vacuum to form a dried polypeptide-PL admixture. Water was then admixed with each dried admixture in an amount calculated to equal 90% of the volume necessary to give a final PL concentration of either 4 or 10 mg/ml

20 (as indicated in Table 7) to form a second admixture. This second admixture was maintained for one hour at 43°C with agitation. Subsequently, a volume of 6% NaCl equal to 10% of the volume necessary to give the desired PL concentration was admixed with the second 25 admixture and the resulting final admixture was maintained for 10 minutes at 43°C. In most cases, the final admixture was subjected to a last step of 3 cycles of freezing and thawing.

B. Results

30 The synthetic surfactants illustrated in Table 6 were prepared as indicated in the table.

TABLE 6

	Peptide	Solvent	(1) Admixture Formed suspension	(2) Phos- pholipid Admixture Method		(3) Composite Hydro- phobicity Value
				A	B	
5	p1-15	n-propyl alcohol	suspension	A	-16.7	
	p11-25	H ₂ O	solution	B	+ 1.7	
	p21-35	Chloroform	suspension	A	- 9.2	
10	p31-45	H ₂ O	solution	B	- 9.9	
	p41-55	H ₂ O	solution	B	- 5.4	
	p51-65	H ₂ O	suspension	B	- 2.2	
	p61-75	methanol	suspension	A	- 9.9	
	p71-81	H ₂ O	suspension	B	+ 3.9	
15	p74-81	H ₂ O	solution	B	+ 3.7	
	p66-81	methanol:H ₂ O	suspension	A	- 1.0	
	p52-81	methanol:H ₂ O	suspension	A	- 6.2	

(1) Each polypeptide was admixed with the indicated solvent to achieve a concentration of 2.5 ug of peptide per ul of solvent.

(2) The letters indicate the synthetic surfactant preparation method used. Those methods are described above.

(3) The composite hydrophobicity value of each peptide was determined as described above.

Each of the synthetic surfactants indicated in Table 6 were assayed for surfactant activity as evidenced by their ability to reduce surface tension in vitro using the "bubble assay" of Enhornung as described above.

The results of this study, shown in Figure 6, indicate that a subject polypeptide, when admixed with pharmaceutically acceptable phospholipids, forms a synthetic pulmonary surfactant that has greater surfactant activity than the phospholipids alone, as evidenced by the lower ΔP values. Typically 10% to 80% lower ΔP values were obtained using the polypeptides. It should be noted that the 8 amino acid residue control peptide p74-81, which does not conform to the

teachings of the present invention, did not form a synthetic PS having a greater activity than the phospholipid alone, thus indicating that amino acid residue length is a critical feature.

5 The surfactant activity of additional exemplary polypeptides of this invention was studied using the "bubble assay" as described above. The results of the study are illustrated below in Table 7.

10 Each polypeptide was admixed with the indicated solvent at a concentration of 2.5 mg of polypeptide per ml of solvent. The resulting admixture was observed to determine whether a solution or a suspension of insoluble polypeptide was formed. Those admixtures forming a suspension were further admixed 15 by water bath sonication for 10 seconds to form a very fine suspension, sufficient for pipetting using glass pipettes.

20 After admixture with solvent, each peptide was admixed with phospholipids (PL), DPPC:PG, 3:1, in chloroform in a glass tube so that the amount of polypeptide added equaled one-tenth (10% by weight) of the amount of PL added, to form a synthetic surfactant according to either method A, B or C.

25 Each of the synthetic surfactants was then assayed for surfactant activity as evidenced by their ability to reduce surface tension in vitro in the bubble assay performed as described above. The pressure gradient (ΔP) is a measure of surfactant activity in the polypeptide-PL final admixture which 30 was determined using an Enhorning Surfactometer as described above. Measurements were obtained at time points of 15 seconds (15"), 1 minute (1') and 5 minutes (5') and are expressed as a mean of three independent measurements of the indicated polypeptide-35 PL admixture. Pressure gradient measurements for

comparable samples of PL alone (PL) and natural human surfactants were determined as controls.

The results of this study are shown in Table 7.

TABLE 7

5	Peptide	Solvent	(1) Admixture Formed	(2) Phos- pholipid Admixture Method	(3) Conc. of PL mg/ml	(4) Pressure Gradient		
						15°	3°	5°
10	p1-15	N-propanol	suspension	A	4	0.94	0.82	0.48
	p36-81	50% chloroform 50% methanol	suspension	C+	10	0.90	0.87	0.79
15	p44-80	68% chloroform 32% methanol	solution	C-	10 ⁴	0.77	0.64	0.24
	p46-76	64% chloroform 36% methanol	solution	C+	10	0.90	0.80	0.62
20	p51-72	75% chloroform 25% methanol	suspension	C+	10	1.15	0.76	0.33
	p51-76	37% chloroform 63% methanol	solution	C+	10	0.99	0.91	0.42
25	p51-80	45% chloroform 55% methanol	solution	C+	10	0.92	0.89	0.48
	p51-81	50% chloroform 50% methanol	suspension	C+	10	0.94	0.86	0.64
	p52-81	67% DMF 33% chloroform	solution	A	4	1.33	1.19	0.96
35	p54-72	76% chloroform 24% methanol	suspension	C+	10	1.28	0.92	0.38
	p54-76	71% chloroform 24% methanol	suspension	C+	10	0.92	0.82	0.23
40	p59-80	32% chloroform 68% methanol	suspension	C-	10 ⁴	1.10	0.96	0.64
	p59-81	68% chloroform 32% methanol	solution	C-	4	1.08	1.02	0.75
	p64-80	48% chloroform 52% methanol	solution	C-	10 ⁴	1.07	0.87	0.11
50	p66-81	40% DMF 60% chloroform	suspension	A	4	1.22	1.11	0.84
	p74-81	water	solution	B	4	2.37	2.32	2.31
55	DL4 (31 mer)	47% chloroform 53% methanol	solution	C-	4	2.00	1.80	1.30
	RL4	32% chloroform 68% methanol	solution	C-	4	0.58	0.65	0.33
60	RL8	19% chloroform 81% methanol	suspension	C+	10	0.68	0.69	0.19
	RRL7	49% chloroform 51% methanol	solution	C-	4	1.65	1.25	1.00
	RCL-1	79% chloroform 21% methanol	suspension	C+	10	0.50	0.59	0.06
70	RCL-2	67% chloroform 33% methanol	suspension	C+	10	0.00	0.00	0.00

TABLE 7 - continued

5	Peptide	Solvent	(1) Admixture Formed	(2) Phos- pholipid Admixture Method		(3) Conc. of PL mg/ml	(4) Pressure Gradient		
				C+	C+		15"	1'	5'
10	RCL-3	75% chloroform 25% methanol	suspension	C+		10	0.55	0.51	0.33
	PL				C+	10	>2.50	>2.50	2.33
15	Natural Human Surfactant					10	1.06	0.89	0.79

(1) Whether the initial admixture of peptide and was a solution or a suspension is indicated.

(2) The letters indicate the synthetic surfactant preparation method used. Those methods are described above. A "+" indicates that the final admixture was subjected to a last step of 3 cycles of freezing and thawing. A "--" indicates the step was not performed.

(3) Concentration ("Conc.") of phospholipid (PL) in the final admixture is indicated in milligrams PL per milliliter admixture (mg/ml).

(4) The pressure gradient is a measure of surfactant activity in the polypeptide-PL final admixture as determined using an Enhorning Surfactometer as described in Example 2. Measurements were obtained at three points of 15 seconds (15"), 1 minute (1') and 5 minutes (5') and are expressed as a mean of 3 independent measurements of the indicated polypeptide-PL admixture. Pressure gradient measurements for comparable samples of PL alone (PL) and natural human surfactant are also shown.

(5) These solutions were made at a concentration of 20 mg/ml PL and were diluted to 10 mg/ml prior to testing.

These results indicate that a subject polypeptide, when admixed with pharmaceutically acceptable phospholipids, forms a synthetic pulmonary surfactant that has a greater surfactant activity than the

phospholipids alone, as demonstrated by the higher volume per given pressure.

Example 3 - In Vivo Assessment of Synthetic Surfactant

5 Activity

A. Methods

Preparation of Synthetic Surfactants

A subject polypeptide was first admixed with solvent as described in Example 2. The resulting

10 admixture was further admixed with phospholipid (PL) so that the amount of polypeptide added was either 3%, 7% or 10% by weight of the amount of PL added as indicated below. The final polypeptide, PL admixture (synthetic surfactant) was formed according to method

15 C using the final freeze thaw step as described in detail in the "Preparation of Synthetic Surfactants" section in Example 2, except that the final admixture had a concentration of 20 mg phospholipid per ml of final admixture.

20

Instillation Protocols

Protocol 1: Fetal rabbits were treated by injection into the trachea of a 0.1 ml solution that contained either a synthetic surfactant prepared in

25 Example 3A or either 2 mg of native surfactant (NS) prepared as described in Example 1 or 2 mg PL.

Protocol 2: Synthetic surfactant was instilled in rabbit fetal lung by injection into the trachea from a single syringe of the following three

30 components such that the components exit the syringe in the following order: (1) 0.05 ml air; (2) 0.1 ml of a synthetic surfactant prepared in Example 3A or either 2 mg of PL or 2 mg of native surfactant; and (3) 0.1 ml air.

Protocol 3: From one syringe, a 0.1 ml aliquot of synthetic surfactant prepared as described in Example 3A (or 2 mg of NS or of PL), was instilled into the rabbit trachea as described above, followed by injection of 0.05 ml lactated Ringer's Solution and 0.2 ml air from a second syringe.

Protocol 4: From one syringe, 0.1 ml of a synthetic surfactant prepared as described in Example 3A (or 2 mg of NS or of PL), 0.15 ml air, 0.1 ml saline, and 0.3 ml air were injected into the trachea as described above. Two subsequent aliquots of 0.3 ml air were given.

Protocol 5: Fetal rabbits were treated by injection into the trachea from a single syringe the following four components such that the four components exit the syringe upon injection in the order listed: (1) 0.2 ml solution that contains either a synthetic surfactant prepared in Example 3A or either 4 mg of native surfactant, or 4 mg PL; (2) a 0.15 ml volume of air; (3) a 0.1 ml normal saline solution; and (4) a 0.3 ml volume of air. The above injection was then repeated 15 minutes after the first injection.

Protocol 6 Rabbits were treated as described in Protocol 5, except that two subsequent aliquots of 0.3 ml air were given following the initial instillation and no additional instillation was given at 15 min.

Fetal Rabbit Model for Studying Surfactant Activity

The surfactant activity of exemplary polypeptides of this invention was studied using the methods described in detail previously by Revak, et al, Am. Rev. Respir. Dis., 134:1258-1265 (1986), with the exceptions noted hereinbelow.

Twenty-seven day gestation fetal rabbits were delivered by hysterotomy and immediately injected with 0.05 ml Norcuron (Organon, Inc., NJ) to prevent spontaneous breathing. The fetal rabbits were then

5 weighed and a small cannula was inserted into the trachea by tracheotomy. Synthetic surfactant prepared as described above was then instilled into the fetal rabbit lung by one of the above instillation protocols.

10 Following instillation the rabbit was placed in a specially designed plethysmograph (containing a Celeesco transducer) connected to a ventilator (Baby Bird, Baby Bird Corp., Palm Springs, CA) and the instilled lung was ventilated at a rate of 30 cycles

15 per minute with a peak inspiratory pressure of 25 cm H₂O, a positive end expiratory pressure of 4 cm H₂O and an inspiratory time of 0.5 seconds. In some studies, dynamic compliance measurements were made at various times throughout the ventilation procedure. In

20 others, static compliance measurements were made following ventilation.

Static compliance measurements were made after 30 minutes of ventilation. The animals were removed from the ventilator and the lungs were degassed at -20 cm H₂O in a bell jar under vacuum. Thereafter, the lungs were first inflated and then deflated through a T-connector attached to a tracheostomy tube. The volume of air required to reach static pressures of 5, 10, 15, 20, 25 and 30 cm H₂O was measured during both

25 inflation and deflation phases to generate static pressure/volume curves as a measure of static compliance.

Using the plethysmograph, dynamic compliance measurements were made at various times throughout a

30 60 minute ventilation period. Computer-assisted data

analysis resulted in compliance data expressed as ml of air per cm H₂O per gram of body weight at each time point. Compliance was calculated by the formula below.

5

$$\text{Compliance} = \frac{\Delta V}{\Delta P}$$

10 ΔP_{tp} = $(C)^{-1} \cdot (\Delta V) + (R) \cdot (F)$

15 P_{tp} = transpulmonary pressure

20 C = compliance (elastic component - relates change in volume to pressure)

25 R = resistance (relates flow to pressure)

30 F = flow

35 V = volume = the integral of flow with respect to time

The above equation was solved with a multiple linear regression for C and R. The compliance (C) represents the elastic nature of the lungs and the resistance (R) represents the pressure necessary to overcome the resistance to the flow of gas into and out of the lungs.

40

B. Results

Static compliance data using instillation protocols 1 and 5 are shown in Figures 7 and 8, respectively. Improved lung compliance was seen in all lungs treated with natural surfactant or with the synthetic surfactants tested as compared with those lungs treated with phospholipids (PL) alone, with one exception. The synthetic surfactant prepared using pl-15 (Figure 7) did not produce improved lung compliance over PL alone when measured by static compliance.

The results of the dynamic compliance studies are illustrated in Table 8.

TABLE 8

5	Peptide Compared To PL	Dynamic Compliance In ml air/cm H ₂ O x 10 ³ g body weight						Sample Given By Protocol #	
		Minutes after Surfactant Instillation							
		10	20	30	40	50	60		
10	PL	7	8	7	10	11	15	4	
		24	22	23	23	22	20	4	
		15	16	17	18	21	29	4	
15	NS	265	251	168	186	173	147*	4	
		418	388	405	288	237	*	4	
		155	176	172		172	179	4	
20	P36-81	5%		255			146*	3	
		5%		245			291	3	
		10%		154			1,162	2	
		10%		252			623	2	
25	P44-80	10% ¹	27	87	138	207	323	6	
		10% ¹	20	23	35	59	87	6	
30	P51-80	10% ¹	42	114	247	300		6	
		10% ¹						6	
35	P52-81	5%		517			226*	3	
		5%		434			55*	3	
		10%		195			1,243	2	
		10%		43			1,690	2	
40	P51-76	10%	33	22	56	87	124	4	
		10%	10	11	186	358	141	4	
		10%	15	36	109	241	264	4	
45	P51-80	10%	17	41	52	78	99	4	
		10%	76	94	149	149	217	4	
		10%	23	71	130	156	182	4	
		10%	42	114	247	388	109*	6	
50	P59-80	10% ¹	24	34	68	107	146	6	
		10% ¹	55	111	190	300	376	6	
55	P64-80	10% ¹	63	129	235	318		6	

1 Prior to instillation into the rabbits, these samples were filtered through a 25 micron filter.

60 * A decrease in compliance with time may indicate the development of pneumothorax.

65 As shown in Table 8, each of the synthetic surfactants of this invention and natural surfactant

improved dynamic compliance values in comparison to phospholipid alone.

C. Discussion

5 The in vivo compliance studies demonstrate that the use of a number of exemplary synthetic surfactants of this invention resulted in enhanced compliance in comparison to phospholipid alone for each of the assayed synthetic surfactants. Thus, the 10 proteins and polypeptides of this invention when admixed with pharmaceutically acceptable phospholipids form synthetic surfactants that have greater surfactant activity than phospholipid alone. Use of the synthetic surfactants is advantageous in producing 15 improved compliance values in vivo.

Example 4 - Study of Binding of C-Terminal Peptide to Lung Epithelial Cells

A. Methods

20 Peptide Binding Assay

A peptide having residues 74-80 of SP18 (VLRCSMD) was radiolabeled by the Bolton-Hunter method (Bolton et al, Biochem J., 133:529-538 1973) with ¹²⁵I (New England Nuclear - 34.1 moles/ml, 28.0 ng/ml, 75 25 µCi/ml).

Human pulmonary epithelial cells (human lung carcinoma cell, ATCC reference no. CCL 185, commonly known as A549 cells) were grown to confluence in 6 well tissue culture dishes. The following solutions 30 were used in this study:

PBS/BSA:

10 mM Na Phosphate + .15 M NaCl + 0.5% BSA pH 7.4

35 Lysis Buffer:

1% SDS in water

Solution F:

5 ml PBS/BSA + 51.56 µg cold peptide

5 Solution D: 2.5 ml PBS/BSA + 87 μ l 125 I-peptide
 5 Solution D1/5: 0.5 ml D + 2.0 ml PBS/BSA
 10 Solution D/125: 0.5 ml D1/5 + 2.0 ml PBS/BSA
 10 Solution E: 2.5 ml PBS/BSA + 87 μ l 125 I-peptide
 10 + 20.78 μ g cold peptide
 10 Solution E1/5: 0.5 ml E + 2.0 ml PBS/BSA
 15 Solution E 1/25: 0.5 ml E1/5 + 2.0 ml PBS/BSA
 15
 20 Three 6 well plates were pretreated by incubating
 20 with 0.5 ml of the following solutions for 15 min. at
 22°C. The odd-numbered wells were pretreated with
 20 PBS/BSA and the even-numbered wells with solution F.
 20 Following removal of the pretreatment solution, the
 20 wells were incubated with 0.5 ml of the following
 20 solutions at 22°C. for the indicated time while gently
 20 rocking the plates.

	<u>Well</u>	<u>Sample</u>	<u>Incubation Time</u>
30	1	D	7 minutes
	2	E	7 minutes
	3	D 1/5	7 minutes
	4	E 1/5	7 minutes
	5	D 1/25	7 minutes
	6	E 1/25	7 minutes
35	7	D	30 minutes
	8	E	30 minutes
	9	D 1/5	30 minutes
	10	E 1/5	30 minutes
	11	D 1/25	30 minutes
	12	E 1/25	30 minutes
40	13	D	143 minutes
	14	E	143 minutes
	15	D 1/5	143 minutes
	16	E 1/5	143 minutes
	17	D 1/25	143 minutes
	18	E 1/25	143 minutes

At the end of the incubation time the supernatant was removed from each well and saved for counting. Each well was washed four times with cold (4°C.) PBS/BSA. The washes were saved for counting. The plate was then brought back to room temperature and 1 ml of lysis buffer was added to each well. The plate was gently shaken until all cells had lysed and came off the plate (3-4 minutes). The solution was removed from each well and counted. A second ml of lysis buffer was added to each well, mixed a few minutes and removed for counting of bound counts. The percent and absolute amounts of counts bound were determined.

Specific counts bound were determined by subtracting the counts bound in wells containing unlabeled (cold) peptide from the corresponding well without cold peptide. The results are illustrated in Table 9, below.

The procedure was repeated with the following changes:

D₁ = 1433 μ l PBS/BSA + 167 μ l ¹²⁵I-peptide [1.78 pmol/500 μ l]

D₂ = 183.3 μ l PBS/BSA + 366.7 μ l D₁ [1.19 pmol/500 μ l]

D₃ = 275 μ l PBS/BSA + 275 μ l D₁ [0.89 pmol/500 μ l]

D₄ = 366.7 μ l PBS/BSA + 183.3 μ l D₁ [0.59 pmol/500 μ l]

D₅ = 458.3 μ l PBS/BSA + 91.7 μ l D₁ [0.30 pmol/500 μ l]

D₆ = 513.3 μ l PBS/BSA + 36.7 μ l D₁ [0.12 pmol/500 μ l]

E₁ = 1386.24 μ l PBS/BSA + 167 μ l ¹²⁵I-peptide [4.676 ng] + 46.76 μ l cold peptide at 100 μ g/ml [4.676 μ g]

E₂-E₆ Diluted as above for D₂ - D₆.

F = 3.398 ml PBS/BSA + 102.29 μ l cold peptide at 100 μ g/ml

Two six-well plates were washed once with 1 ml PBS/BSA. The odd numbered-wells were pretreated with PBS/BSA and the even-numbered wells with solution F. Following removal of the pretreatment solution, the 5 following solutions were added.

	<u>Well</u>	<u>Sample</u>	<u>Well</u>	<u>Sample</u>
10	1	D ₁	7	D ₄
	2	E ₁	8	E ₄
	3	D ₂	9	D ₅
	4	E ₂	10	E ₅
	5	D ₃	11	D ₆
	6	E ₃	12	E ₆

The solutions were incubated for 30 minutes at room temperature with gentle rocking. The 15 supernatants were then removed and saved for counting. Each well was washed 4 times with 0.5 ml of cold PBS/BSA. Washes were saved for counting. 1 ml of 1% SDS was added to each well to solubilize the cells. After 3 minutes all the cells could be seen to have 20 come off the plate. The lysed cell-containing supernatant was counted, together with a second SDS wash of the wells. Total counts and the percentage of counts bound were determined. Specific binding was determined by subtracting the counts bound in wells 25 containing cold peptide from the corresponding well without cold peptide. The results are illustrated in Table 10.

B. Results

30 The results of the binding studies are illustrated below in Tables 9 and 10.

TABLE 9

5	Well	Total CPM	Total CPM Bound	% Bound	Difference	Specific Cts Bound
10	1	1,109,126	24,414	2.23	0.63%	6,930
	2	1,087,659	17,353	1.60		
15	3	223,170	4,479	2.01	0.15%	330
	4	221,608	4,113	1.86		
20	5	45,877	828	1.80	-0.04%	- 18
	6	47,731	880	1.84		
25	7	1,103,606	25,905	2.35	0.68%	7,480
	8	1,152,287	19,230	1.67		
30	9	227,396	4,996	2.20	0.41%	901
	10	230,974	4,137	1.79		
35	11	47,899	1,030	2.15	0.30%	132
	12	49,894	922	1.85		
40	13	1,151,347	10,071	.87	0.01%	110
	14	1,108,755	9,506	.86		
45	15	220,340	1,692	.77	-0.02%	- 44
	16	229,253	1,800	.79		
50	17	46,893	407	.87	0.06%	26
	18	47,426	386	.81		

35 * Corrected to 1,100,000 cpm/undiluted tube.

TABLE 10

40	Well	Total CPM	CPM Bound	% Bound	Corrected Difference	p mol
45	1	3,070,705	66,954	2.78	9,390	1.78
	2	2,995,775	56,055	1.87		
50	3	2,029,323	39,562	1.95	5,723	1.19
	4	2,013,557	33,573	1.67		
55	5	1,436,189	26,883	1.87	1,755	.89
	6	1,427,731	25,073	1.76		
60	7	994,288	15,669	1.58	503	.59
	8	964,481	14,776	1.53		
65	9	460,317	6,513	1.41	- 52	.30
	10	479,746	6,816	1.42		
70	11	202,494	2,930	1.45	0	.12
	12	192,990	2,806	1.45		

* Corrected for 1.78 p mol = 3,033,740 cpm

C. Discussion

As can be seen from the data in Table 9, the study demonstrated that the peptide was binding

specifically to the cells as demonstrated by competitive inhibition by unlabeled peptide. However, the cells were not saturated by the amount of labeled peptide used in this study. Additionally, degradation 5 of the peptide was occurring by 143 minutes.

The second study was performed using the 30 minute incubation period and an increased amount of labeled peptide to achieve saturation of the cells. As seen in Table 10, specific binding was again demonstrated. 10 Further, saturation was achieved as demonstrated by leveling-off of the amount of bound counts at high concentration of labeled peptide.

The binding studies thus demonstrate that the C-terminal peptide of this invention binds 15 specifically to lung epithelial cells.

The foregoing specification, including the specific embodiments and examples, is intended to be illustrative of the present invention and is not to be 20 taken as limiting. Numerous other variations and modifications can be effected without departing from the true spirit and scope of the present invention.

WHAT IS CLAIMED IS:

1. A composition comprising human SP18 monomer and human SP18 dimer that is free of other pulmonary surfactant proteins.

5

2. The composition of claim 1 wherein said proteins are admixed with a pharmaceutically acceptable phospholipid to form a synthetic pulmonary surfactant.

10

3. A composition comprising substantially pure human SP18 monomer protein.

4. The composition of claim 3 wherein said protein is admixed with a pharmaceutically acceptable phospholipid to form a synthetic pulmonary surfactant.

15 5. A DNA segment consisting essentially of a DNA sequence encoding mature human SP18 monomer protein.

20

6. The DNA segment of claim 5 wherein said sequence corresponds to the DNA sequence shown in Figure 1 from nucleotide position 187 to nucleotide position 429.

25

7. A recombinant nucleic acid molecule comprising a vector operatively linked to a structural gene capable of expressing, without post-translational proteolytic modification, mature human SP18 monomer protein.

30 8. The recombinant nucleic acid molecule of claim 7 wherein said structural gene is operatively linked to an expression control element.

35

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2. The composition of claim 1 wherein said proteins are admixed with a pharmaceutically acceptable phospholipid to form a synthetic pulmonary surfactant.

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4. The composition of claim 3 wherein said protein is admixed with a pharmaceutically acceptable phospholipid to form a synthetic pulmonary surfactant.

20

5. A DNA segment consisting essentially of a DNA sequence encoding mature human SP18 monomer protein.

25

6. The DNA segment of claim 5 wherein said sequence corresponds to the DNA sequence shown in Figure 1 from nucleotide position 187 to nucleotide position 429.

25

7. A recombinant nucleic acid molecule comprising a vector operatively linked to a structural gene capable of expressing, without post-translational proteolytic modification, mature human SP18 monomer protein.

35

8. The recombinant nucleic acid molecule of claim 7 wherein said structural gene is operatively linked to an expression control element.

9. A method of preparing human SP18 monomer protein comprising:

(a) initiating a culture, in a nutrient medium, of mammalian cells transformed with a recombinant DNA molecule comprising a vector operatively linked to a structural gene capable of expressing, without post-translational proteolytic modification, human SP18 monomer;

5 10 (b) maintaining said culture for a time period sufficient for said cells to express human SP18 monomer protein from said recombinant DNA molecule; and

15 (c) recovering said protein from said culture.

10 15 10. A polypeptide consisting essentially of at least 10 amino acid residues and no more than about 60 amino acid residues, said polypeptide corresponding in sequence to an amino acid residue sequence of human SP18 monomer, said polypeptide, when admixed with a 20 pharmaceutically acceptable phospholipid forming a synthetic pulmonary surfactant having a surfactant activity greater than the surfactant activity of the phospholipid alone.

25 11. The polypeptide of claim 10 further characterized as having at least one sequence corresponding to at least 15 contiguous amino acid residues in the linear sequence of SP18 located from residues 1 to 75.

30 35 12. The polypeptide of claim 11 having an amino acid residue sequence selected from the group consisting of:

FPIPLPYCWLCRALI,
CRALIKRIQAMIPKG,

MIPKGALAVAVAQVC,
VAQVCRVVPPLVAGGI,
VAGGICQCLAERYSV,
CQCLAERYSVILLDTLLGRMLPQLVCRLVLR
5 ERYSVILLDTLLGRM,
ERYSVILLDTLLGRMLPQLVCR,
ERYSVILLDTLLGRMLPQLVCRLVLR,
SVILLDTLLGRMLPQLVCR,
SVILLDTLLGRMLPQLVCRLVLR, and
10 LLGRMLPQLVCRLVLR.

13. The polypeptide of claim 10 further characterized as having a carboxy-terminal sequence corresponding to at least 5 contiguous amino acid residues in the linear sequence of SP18 and including residue 80.

14. The polypeptide of claim 13 further characterized as additionally having at least one sequence corresponding to at least 15 contiguous amino acid residues in the linear sequence of SP18 located from residues 1 to 75.

15. The polypeptide of claim 13 further characterized as having a carboxy-terminal amino acid residue sequence represented by the formula:

-RLVLRCMDD_Z,

wherein Z is an integer having a value of 0 or 1, such that when Z is 0 the D residue to which it is a subscript is absent and when Z is 1 the D residue to which it is a subscript is present.

16. The polypeptide of claim 15 having an amino acid residue sequence selected from the group consisting of:

CRLVLRCSMDD;
LPQLVCRVLRLRCSMDD;
DTLLGRMLPQLVCRVLRLRCSMDD;
RYSVILLDTLLGRMLPQLVCRVLRLRCSMDD;
5 ERYSVILLDTLLGRMLPQLVCRVLRLRCSMDD;
ERYSVILLDTLLGRMLPQLVCRVLRLRCSMDD;
RVVPLVAGGICQCLAERYSVILLDTLLGRMLPQLVCRVLRLRCSMDD; and
AQVCRVVPLVAGGICQCLAERYSVILLDTLLGRMLPQLVCRVLRLRCSMDD.

10 17. A polypeptide consisting essentially of at least 10 amino acid residues and no more than about 60 amino acid residues having alternating hydrophobic and hydrophilic amino acid residue regions represented by the formula $(Z_aU_b)_cZ_d$, wherein:

15 Z is a hydrophilic amino acid residue independently selected from the group consisting of R, D, E and K;
U is a hydrophobic amino acid residue independently selected from the group
20 consisting of V, I, L, C, Y and F;
a has an average value of about 1 to about 5;
b has an average value of about 3 to about 20;
c is 1 to 10; and
d is 1 to 3;

25 said polypeptide, when admixed with a pharmaceutically acceptable phospholipid, forming a synthetic pulmonary surfactant having a surfactant activity greater than the surfactant activity of the phospholipid alone.

30 18. The polypeptide of claim 17 having an amino acid residue sequence selected from the group consisting of:

35 DLLLDLRLDLDLRLDLDLRLD,
RLLLRLLRLRLRLRLRLR,
RLLLLLRLRLRLRLRLR,

RLLLLLRLRPLLLLLLRL,
RLLLLCLLRLLLLCLLR,
RLLLLCLLRLLLLCLLRLL, and
RLLLLCLLRLLLLCLLRLLLLCLLR.

5

19. A composite polypeptide of at least 10 and no more than 60 amino acid residues consisting essentially of an amino terminal sequence and a carboxy terminal sequence wherein:

10 said amino terminal sequence has at least 10 amino acid residues and has a composite hydrophobicity of less than 0; and

15 said carboxy terminal sequence is characterized as consisting essentially of an amino acid residue

15 sequence represented by the formula:

-RLVLRCSMDD_z,

wherein Z is an integer having a value of 0 or 1, such that when Z is 0 the D residue to which it is a subscript is absent and when Z is 1 the D residue to 20 which it is a subscript is present;

25 said composite polypeptide, when admixed with a pharmaceutically acceptable phospholipid forming a synthetic pulmonary surfactant having a surfactant activity greater than the surfactant activity of the phospholipid alone.

20. The composite polypeptide of claim 19 wherein said amino terminal sequence is further characterized as having alternating hydrophobic and hydrophilic 30 amino acid residue regions represented by the formula $(Z_a U_b)_c Z_d$, wherein:

Z is a hydrophilic amino acid residue independently selected from the group consisting of R, D, E and K;

35